PDB annotation					COMPLEX (IMMUNOGLOBULIN/LIPOPROTEIN) OSPA; COMPLEX (IMMUNOGLOBULIN/LIPOPROTEIN), OUTER SURFACE 2 PROTEIN A COMPLEXED WITH FABI84.1, BORRELIA BURGDORFERI 3 STRAIN B31			CELL ADHESION PROTEIN NCAM MODULE 2, CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	LIGASE CYCLIN A/CDK2. ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRK, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE
Coumpound	GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	TLYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) IHNG 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) 1HNG 3	FAB 184.1; CHAIN: L, H; OUTER SURFACE PROTEIN A; CHAIN: O;	MUSCLE PROTEIN TITIN MODULE MS (CONNECTIN) ITNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) ITNM 4 ITNM 58	IMMUNOGLOBULIN IGG2A FAB FRAGMENT (CN1206) 2GFB 3	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;	SKP2; CHAIN: A, C, E, G, 1, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;
SeqFold score			60.69		55.92		53.4		
PMF score		0.76		0.35		-0.18		-0.18	0.92
Verify		60.0		0.35		0.09		0	-0.07
PSI- BLAST		9.10E-18	2.60E-18	2.60E-18	0.0011	3.60E-14	3.60E-05	1.80E-09	2.60E-05
End AA		199	217	210	230	213	230	84	81
Start		28	25	53	19	131	61	22	46
Chain ID			V	∢	يا		⋖	<	V V
PDB ID		lhnf	lhng	lhng	losp	Itum Tum	2gfb	3лс п	Ifqv
SEQ ID NO:		741	741	741	741	741	741	741	747

PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) UPSTREAM STIMULATORY FACTOR 1; USF, DNA BINDING, BASIC-HELIX-LOOP-HELIX, LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)		A- COMPLEX (TRANSCRIPTION FACTOR/DNA) NF-KB P50, COMPLEX (TRANSCRIPTION FACTOR/DNA)	Z.	HYDROLASE MALTOGENIC ALPHA AMYLASE; AMYLASE, GLYCOSIDE HYDROLASE, STARCH DEGRADATION		COMPLEX (ZINC FINGER/DNA) EX COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN C;	COMPLEX (ZINC FINGER/DNA) LEX COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN C;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	USF; CHAIN: A, B; DNA; CHAIN: C, D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NUCLEAR FACTOR KAPPA-B; CHAIN: A, B; KB SITE, DNA (5'-D/TOAGAATTCCC)-3'); CHAIN: C, D;	GLYCOSYLTRANSFERASE CYCLODEXTRIN GLUCANOTRANSFERASE (E.C.2.4.1.19) (CGTASE) 1CYG	ALPHA-AMYLASE; CHAIN: A;	CHROMOSOMAL PROTEIN UBIQUITIN IUBI 3	QGSR ZINC FINGER PEPTIDE: CHAIN: A: DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE, CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold									
PMF	60.0	0.41	0.04	0.05	0.37	0.03	0.03	0.03	0.04
Verify	-0.61	0.39	-0.37	-0.02	0.11	-0.58	-0.45	-0.33	-0.52
PSI- BLAST	0.0078	0.0013	0.0013	0.0016	0.0061	0.0065	3.60E-25	3.60E-24	1.40E-44
End	390	346	319	1059	967	1617	314	162	220
Start	353	243	234	932	854	1529	229	78	134
Chain ID	∢	<	V		\ <		∢	∢	O
PDB ID	lan4	l ikn	lnfk	lcyg	1qho	lubi	lalh	laih	1me y
SEQ ID NO:	750	750	750	754	754	754	756	756	756

PDB annotation	(ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA INTERACTION PROTEIN DESIGN 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRISIAL SIRUCIURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION. PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	CRYSTAL STRUCTURE COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	INTER ACTION PROTEIN DESIGN 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)
Coumpound	DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E,	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN: C F G:			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA CHAIN A B D F	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		•	DNA, CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	TROTEIN, CHAIN. C. F. G.		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	PROTEIN: CHAIN: D F D.	1 NO 1 LIM, CITAIN: C. 1, C.	
SeqFold score																																	
PMF	90.0			0.28				0.12				-0.07				0.17					0.05				0.07					-0.19			
Verify score	9.0-			-0.15				-0.19				90.0				-0 44					90:0				-0.3				;	90.0			
PSI- BLAST	9.00E-44			3.60E-42				1.10E-42				3.60E-30				1.80F-42	!				5.40E-12				9.00E-11					9.00E-11			
End	284			465				612				627				162					190				284					465		_	
Start	193			379				526				555				77					163				254					437			
Chain 1D	C			ပ				C				U				J					U				G					יב			
PDB ID	Ime	>		Ime	>			lmc	<u>~</u>			Jmc 1	>			Ime	· >			7	lmc :	<u> </u>			Ime	^			1.	e E	`		
SEQ ID NO:	756			952				756				756				756		_			756				756				,	95/		_	

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN. DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION
Coumpound	TRANSCRIPTION FACTOR CON IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F; TRA TRA GEN GEN	TFIIIA; CHAIN: A, D; 5S COP RIBOSOMAL RNA GENE; REC CHAIN: B, C, E, F; (TR REC	TFIIIA; CHAIN: A, D; 5S COI RIBOSOMAL RNA GENE; REC CHAIN: B, C, E, F; (TR REC	YY1; CHAIN: C; ADENO-COI ASSOCIATED VIRUS PS REC INITIATOR ELEMENT DNA; TRA CHAIN: A, B; INI FIN REC	YYI; CHAIN: C; ADENO-CO ASSOCIATED VIRUS P5 REG INITIATOR ELEMENT DNA; TR, CHAIN: A, B; INI FIN REG	YY1; CHAIN: C; ADENO-CO ASSOCIATED VIRUS P5 REG INITIATOR ELEMENT DNA; TR, CHAIN: A, B; NIFEN	YY1; CHAIN: C; ADENO- CO
SeqFold score		69.84					
PMF	0.34		0.11	0.11	0.23	0	0.01
Verify score	-0.43		-0.18	-0.77	-0.21	-0.72	-0.53
PSI- BLAST	1.80E-15	1.40E-34	1.40E-34	1.30E-30	1.80E-27	1.80E-31	1.80E-30
End AA	432	285	302	253	436	578	130
Start AA	349	106	135	139	325	470	85
Chain TD	∢	∢	Y	ပ	U	ပ	၁
PDB ID	1113	1tf6	1116	Iubd	lubd	lubd	lubd
SEQ ID NO:	756	756	756	756	756	756	756

SEQ ID NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
l									ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
										RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
	2adr	 	349	406	3.60E-15	-0.63	0.11		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, A DDI 2 ZNC ENGED MAD
756	2adr		498	554	1.40E-15	-0.13	0.04		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION ARANSCRIPTION REGULATION ARANSCRIPTION REGULATION, ABRI. ZINC FINGER, NMR.
756	2gli	<	142	286	1.10E-28	-0.53	0.07		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
	Icun	<	537	726	7.80E-14	0.12	-0.02		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, TANDEM 1-HELIX COILED-COILS,
	lcun	∢	999	749	9.10E-15	0.13	0		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
	orp1		1096	1203	5.20E-06	80.0	0.43		BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7	CYTOSKELETON
	1fb8	¥	1093	1203	5.20E-18	0.44	0.89		DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN, ADAPTOR
_	1fgy	A	1097	1203	1.20E-18	0.56	0.68		GRP1; CHAIN: A:	SIGNALING PROTEIN ARF! GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN
	1b0x	A	801	857	3.60E-12	0.04	-0.09		EPHA4 RECEPTOR TYROSINE KINASE; CHAIN:	TRANSFERASE RECEPTOR TYROSINE KINASE, PROTEIN INTERACTION

	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ΒŞ	<u>e</u>	<u>a</u>	¥	AA.	BLAST	score	score	score		
. [A;	MODULE, 2 DIMERIZATION DOMAIN, TRANSFERASE
762	1b4f	¥	793	863	1.10E-14	0.1	-0.02		EPHB2; CHAIN: A, B, C, D, E, F, G, H;	SIGNAL TRANSDUCTION SAM DOMAIN, EPH RECEPTOR, SIGNAL TRANSDUCTION, OLIGOMER
762	lsgg		797	861	7.20E-14	0.53	-0.02		EPHRIN TYPE-B RECEPTOR 2; CHAIN: NULL;	TYROSINE-PROTEIN KINASE NMR, RECEPTOR OLIGOMERIZATION, EPH RECEPTORS, TYROSINE 2 PHOSPHORYLATION, SIGNAL TRANSDUCTION, TYROSINE-PROTEIN 3 KINASE
763	184р	₹	-	92	1.30E-20			69.57	S100A10: CHAIN: A, B;	CALCIUM/PHOSPHOLIPID BINDING PROTEIN P11. CALPACTIN LIGHT CHAIN; S100 FAMILY. EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN
763	1a4p	<	2	68	1.30E-20	0.67			S100A10; CHAIN: A, B;	CALCIUM/PHOSPHOLIPID BINDING PROTEIN P11, CALPACTIN LIGHT CHAIN, S100 FAMILY, EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN
763	lak8		4	28	3.60E-21	0.11	-0.11		CALMODULN; CHAIN: NULL;	CALCIUM-BINDING PROTEIN CALMODULIN CERIUM TRIC- DOMAIN, RESIDUES 1 - 75; CERIUM- LOADED, CALCIUM-BINDING PROTEIN
763	164c	≺	-	92	7.20E-19			145.44	S-100 PROTEIN, BETA CHAIN; CHAIN: A, B;	METAL BINDING PROTEIN S100B, S100BETA; S100BETA, S100B, NMR, DIPOLAR COUPLINGS, EF-HAND, S100 2 PROTEIN, CALCIUM- BINDING PROTEIN, FOUR-HELIX BUNDLE, THREE- 3 DIMENSIONAL STRUCTURE
763	1cb1		3	83	7.80E-27			57.95	CALCIUM-BINDING PROTEIN CALBINDIN D9K (INTACT FORM) (NMR. 13 STRUCTURES) ICBI 3	
763	1cb1		4	82	7.80E-27	0.43	0.99		CALCIUM-BINDING	

PDB Chain Start ID ID AA	<u> </u>	Start		End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									PROTEIN CALBINDIN D9K (INTACT FORM) (NMR, 13 STRUCTURES) ICBI 3	
m 4 81 9	4	18		6	9.00E-21	0.21	0.06		CALCIUM-BINDING PROTEIN CALMODULIN COMPLEXED WITH CALMODULIN-BINDING DOMAIN OF ICDM 3 CALMODULIN-DEPENDENT PROTEIN KINASE II ICDM 4	
lemf 1 81 7				7	7.20E-21	0.17	0.15		CALMODULIN (VERTEBRATE); ICMF 6 CHAIN: NULL; ICMF 7	CALCIUM-BINDING PROTEIN CALMODULIN APO TR2C-DOMAIN; ICMF 9
lexr A 4 89 3	4 89	68	ļ	3	3.60E-23	90.0	-0.17		CALMODULIN; CHAIN: A;	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER
171 A 8 81 7	88	81		_	7.20E-20	-0.17	0.18		CALMODULIN; CHAIN: A;	TRANSPORT PROTEIN CALCIUM BINDING, EF HAND, FOUR-HELIX BUNDLE
lmli 2 89 1.	68	68			1.40E-16			137.02	S-100 PROTEIN; CHAIN: NULL;	CALCIUM-BINDING CALCIUM- BINDING, ZINC, METAL-BINDING, ACETYLATION
11op 5 92 7	92	92		7.	7.20E-20	0.07	-0.07		CONTRACTILE SYSTEM PROTEIN TROPONIN C I TOP 3	
lajs A 135 252 2.	135 252	252		7,	2.60E-09	-0.18	0.09		CALPAIN; CHAIN: A, B;	CALCIUM-BINDING PROTEIN CALCIUM-BINDING PROTEIN, CALCIUM-DEPENDENT PROTEASE, APO 2 FORM, SMALL SUBUNIT
1dtl A 131 252 2	131 252	252	 	7	2.60E-09	-0.29	0.28		CARDIAC TROPONIN C: CHAIN: A;	STRUCTURAL PROTEIN HELIX-TURN- HELIX
lexr A 129 252 1	129 252	252		_	1.30E-09	-0.18	0.39		CALMODULIN; CHAIN: A;	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER
252	129 252	252		7	2.60E-08	0.05	-0.03		CALCIUM-BINDING PROTEIN NCS-1; CHAIN: A;	METAL BINDING PROTEIN YEAST FREQUENIN EF-HAND, CALCIUM
1tcf 135 252 3	252	252		<u> </u>	3.90E-10	-0.12	0.47		TROPONIN C, CHAIN: NULL,	CALCIUM-REGULATED MUSCLE CONTRACTION MUSCLE CONTRACTION, CALCIUM-BINDING, TROPONIN, E-F HAND, 2 OPEN
										CONFORMATION REGULATORY DOMAIN, CALCIUM-REGULATED 3

Coumpound PDB annotation	MUSCLE CONTRACTION	C; ITNX 4 CALCIUM-BINDING PROTEIN EF- L; ITNX 5 HAND ITNX 14	CALCIUM BINDING PROTEIN CALMODULIN (/TR=2=C\$ FRAGMENT COMPRISING RESIDUES 78 - 148 ITRC 3 OF THE INTACT MOLECULE) ITRC 4	CALCIUM-BINDING PROTEIN PARVALBUMIN (ALPHA LINEAGE) SPAL 3	AQUAPORIN-1; CHAIN: A; MEMBRANE PROTEIN AQPI WATER CHANNEL, TWO-DIMENSIONAL CRYSTAL, ELECTRON 2 DIFFRACTION, ELECTRON MICROSCOPY	AQUAPORIN-1: CHAIN: A; MEMBRANE PROTEIN AQP1 WATER CHANNEL, TWO-DIMENSIONAL CRYSTAL, ELECTRON 2 DIFFRACTION, ELECTRON MICROSCOPY	SYNAPTOTAGMIN I; CHAIN: ENDOCYTOSIS/EXOCYTOSIS A; SYNAPTOTAGMIN, C2-DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSIS	.B;	AGMIN III; ENDOCYTOSIS/EXOCYTOSIS BETA SANDWICH, CALCIUM ION, C2 DOMAIN	PROTEIN KINASE C, ALPHA TRANSFERASE CALCIUM++, PHOSPHOLIPID BINDING PROTEIN, CALCIUM-BINDING 2 PROTEIN, PHOSPHATIDYLSERINE, PROTEIN KINASE C	CALCIUM/PHOSPHOLIPID
		TROPONIN C; 1TNX 4 CHAIN: NULL; 1TNX 5	CALCIUM BINDING PROTEIN CALMODULI (/TR=2=C\$ FRAGMENT COMPRISING RESIDUE 148 ITRC 3 OF THE INT MOLECULE) ITRC 4	CALCIUM-BINDING PROTEIN PARVALBI (ALPHA LINEAGE) 5	AQUAPORII	AQUAPORII	SYNAPTOT. A;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN	SYNAPTOTAGMIN III. CHAIN: A;	PROTEIN KINASI TYPE; CHAIN: A;	CALCIUM/P
SeqFold score											
PMF		0.31	0.95	0.53	0.51	0.88	0.94	0.01	0.82	0.71	-
Verify score		-0.53	-0.37	0.27	-0.22	-0.13	0.21	-0.13	0.38	0.48	0.3
PSI- BLAST		1.00E-10	1.30E-08	1.30E-08	1.10E-73	3.90E-76	1.30E-11	5.20E-05	6.50E-07	0.0026	6.50E-12
End		252	252	252	227	227	664	252	647	999	664
Start		140	194	171	01	6	541	26	541	541	541
Chain ID			<		∢	<	<	В	∢	∢	
PDB ID		ltnx	Itro	5pal	1fqy	1fqy	1byn	1dn1	ldqv	Idsy	Irsy
SEQ ID		992	766	766	768	768	769	692	692	769	769

PDB annotation		P21; SOS; COMPLEX (ONCOGENE PROTEIN/EXCHANGE FACTOR), SMALL GTPASE, 2 EXCHANGE FACTOR	P21; SOS; COMPLEX (ONCOGENE PROTEIN/EXCHANGE FACTOR), SMALL GTPASE, 2 EXCHANGE FACTOR	P21; SOS; COMPLEX (ONCOGENE PROTEIN/EXCHANGE FACTOR), SMALL GTPASE, 2 EXCHANGE FACTOR	SIGNAL TRANSDUCTION PROTEIN	CYTOSKELETON		SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN	SIGNALING PROTEIN ARF! GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN	
Coumpound	BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	H-RAS, CHAIN: R. SON OF SEVENLESS-1; CHAIN: S;	H-RAS; CHAIN; R; SON OF SEVENLESS-1; CHAIN: S;	H-RAS; CHAIN: R; SON OF SEVENLESS-1; CHAIN: S;	BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7	SIGNAL TRANSDUCTION PROTEIN DYNAMIN (PLECKSTRIN HOMOLOGY DOMAIN) (DYNPH) 1DYN 3	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	GRP1; CHAIN: A;	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL
SeqFold score		79.72									
PMF			_	_	9.68	0.4	0.25	0.9	0.05	0.36	0.24
Verify score			0.2	0.26	-0.09	-0.22	-0.04	0.3	0.04	0.35	-0.02
PSI- BLAST		9.10E-77	9.10E-77	7.20E-61	1.40E-18	7.20E-12	5.40E-06	1.00E-07	1.30E-12	1.10E-16	1.10E-13
End AA		311	291	312	541	546	541	546	539	546	542
Start		_	12	41	438	439	456	429	441	447	437
Chain ID		S	S	S			V	₹	¥	٧	
PDB 1D		1bkd	15kd	15kd	1 btn	1 dro	ldyn	1168	168	lfgy	lpls
SEQ NO:		770	770	770	170	770	770	770	770	770	770

ind PDB annotation	40L0GY 1T IPLS 3 IIS)6 I(NS(G105- IR, 25 LS 5		ER COMPLEX (ZINC FINGER/DNA) A; DUPLEX COMPLEX (ZINC FINGER/DNA), ZINC DE FINGER, DNA-BINDING PROTEIN HAIN: B, C;	S, D, E, FINGER, PROTEIN-DNA C, F, G, CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA) FINGER, PROTEIN-DNA CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	S, D, E; COMPLEX (ZINC FINGER/DNA) ZINC C FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ER .;	3. D. E. COMPLEX (ZINC FINGER/DNA) ZINC C FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	C FINGER FINGER, PROTEIN-DNA C F. G; NERACTION, PROTEIN DESIGN, 2
Coumpound	PLECKSTRIN HOMOLOGY DOMAIN) MUTANT 1PLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS 1PLS 4 (INS(G105- LEHHIHHHH)) (NMR, 25 STRUCTURES) 1PLS 5	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A, DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA: CHAIN: A. B. D. E. CONSENSUS ZINC FINGER PROTEIN; CHAIN: C. F. G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold			78.23					
PMF		0.99		1-	-	_	_	_
Verify score		0.19		0.37	0.29	0.46	0.29	0.25
PSI- BLAST		6.50E-41	5.20E-45	5.40E-46	3.60E-47	1.80E-48	1.10E-49	3.60E-51
End		210	351	209	237	265	293	377
Start		131	269	128	156	184	212	296
Chain ID		4	Y .	U	O	ပ	ပ	O
PDB ID		lath	lalh	Jme Y	lme y	1me ×	y y	Ime y
SEQ ID		277	772	277	772	772	772	772

PDB Chain	Chai	٥	Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
Ime C	O		296	378	3.60E-51			102.88	DNA; CHAIN: A, B, D, E. CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Ime C	O		324	405	3.60E-51	0.16			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
1me C	၁		352	433	7.20E-51	0.33	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Ime V	ပ		380	461	7.20E-51	0.44	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Ime y	ပ		66	181	5.40E-43	0.04	0.42		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
1tf6 A	<		126	298	5.20E-73			115.72	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN
1tf6 A	<		129	274	1.10E-36	0.22	0.76		TFIJIA; CHAIN: A. D. 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN
Itf6 A	<		297	443	1.30E-37	0.09	0.94		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA

PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX CTB ANGCPIPTION	REGULATION RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGILL ATION DNA) YING-YANG I	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGITATION TO SECURE ATTOMONA) VING-YANG I:	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	PECOCNITION 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1:	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: D C E E:	CIDIN: B, C, E, F,	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA; CHAIN: A. B:			YY1; CHAIN: C; ADENO- ASSOCIATED VIRIIS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS INITIATOR FLEMENT DNA:	CHAIN: A, B;	
SeqFold score							-					_										-	
PMF		96.0		0.15		0.87					_	_			-					_			
Verify score		0.15		-0.27		0.12				0.32					0.2					0.3			
PSI. BLAST		1.80E-36		3.60E-33		3.90E-42				1.80E-32				24 1000	6.50E-52					1.30E-53			
End		461		218		209				237				17.0	/57					566			
Start AA		325		72		120				131				:5:	133					154			
Chain ID		V	,	A	_	၁				ပ				,	ر					၁			
PDB ID		1tf6		1116	:	Iubd				lubd				77.1	D D					lubd			
SEQ ID NO:		772		772		772				277		_		133	7//					772			

PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;	INTIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION PEGII ATTONIONA) YING-YANG I	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYT, ZINC Z	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TING-TANG I;	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(IKANSCRIPTION KEGULATION/DINA)	COMPLEX (TRANSCRIPTION DEGLII ATTON/DNA) VING-YANG 1:	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS PS	CHAIN: A. B.				YYI; CHAIN: C; ADENO-	INITIATOR ELEMENT DNA:	CHAIN: A, B;
SeqFold score		89.18																									
PMF score				0.92				96.0					86.0						_						0.95		
Verify score				-0.04				0.26					0.22						0.16						0.4		
PSI- BLAST		7.80E-55		7.80E-55				6.50E-56					1.60E-35						1.30E-56						5.20E-51		
End		322		349				405					405						434						461		
Start		212		238				294					304						322			~··			350		
Chain ID		U		S)					C						J						ပ		
PDB ID	+	lubd		lubd				lubd					lubd			-			lubd						1 ubd	_	
SEQ ID	ä	772		772				772					772						772						772		

PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BNDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BNDING PROTEIN/DNA)
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII: CHAIN: A; DNA; CHAIN: C. D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score						98.73				
PMF		0.92	0.88	-	0.76		0.93	0.84	96:0	-
Verify		0.32	0.21	0.35	91.0		0.4	0.24	0.35	0.47
PSI- BLAST		1.80E-34	2.60E-54	3.60E-33	3.90E-66	1.00E-71	1.00E-71	7.20E-33	6.50E-67	5.40E-34
End		461	239	264	295	351	407	432	461	460
Start AA		360	121	128	128	212	268	304	324	332
Chain ID		O	<	∀	⋖	∢	<	Ą	¥	¥
PDB ID		1ubd	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli
SEQ ID	į	772	772	772	772	772	772	772	772	772

						7				<u>.</u>	
PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	HYDROLASE TETRATRICOPEPTIDE, TRP, HYDROLASE, PHOSPHATASE, PROTEIN-PROTEIN INTERACTIONS, TPR, 2 SUPER-HELIX, X-RAY STRUCTURE	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN BINDING	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 7 HELICAL REPEAT	AMIDOTRANSFERASE AMIDOTRANSFERASE, THIOESTER	AMIDOTRANSFERASE AMIDOTRANSFERASE, THIOESTER	LYASE AIRC, PURK; ATP-GRASP, CARBOXYPHOSPHATE, PURINE BIOSYNTHESIS, LYASE	LIGASE ATP-GRASP, CARBOXYLASE, BIOTIN-DEPENDENT	LIGASE LMDDLZ; ATP-BINDING.
Coumpound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	SERINE/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;	TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	TPR1-DOMAIN OF HOP; CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTS1- CONTAINING PEPTIDE; CHAIN: C, D;	CARBAMOYL PHOSPHATE SYNTHETASE; CHAIN: A, B, C, D, E, F, G, H;	CARBAMOYL PHOSPHATE SYNTHETASE; CHAIN: A, B, C, D, E, F, G, H;	NS- CARBOXYAMINOIMIDAZOL E RIBONUCLEOTIDE CHAIN: A:	BIOTIN CARBOXYLASE; CHAIN: A, B;	D-ALANINE: D-LACTATE
SeqFold score											
PMF score	-0.05	0.15	0.77	89.0	0.74	66:0	6:0	96.0	-0.11	0.54	0.45
Verify score	0.07	-0.01	60.0	0.14	0.12	0.47	0 14	0.49	0.07	0.32	0.07
PSI- BLAST	9.00E-28	5.40E-31	1.30E-08	9.10E-08	6.50E-10	6.50E-09	1.80E-64	1.10E-39	1.80E-16	1.30E-86	9.00E-37
End	180	211	180	180	180	179	098	892	828	596	858
Start AA	50	71	105	105	105	103	454	529	532	534	531
Chain ID	¥	₹		¥	⋖	<	∢	∀	4	<	A
PDB ID	2gli	2gli	la17	lefr	leiw	1fch	1a9x	1a9x	1b6r	ldv1	Ich:
SEQ ID NO:	772	772	773	773	773	773	774	774	774	774	774

	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
	1chj	В	531	858	1.80E-36	0.22	0.63		D-ALANINE:D-LACTATE LIGASE: CHAIN: A, B;	LIGASE LMDDL2; ATP-BINDING. GRASP MOTIF FOR ATP.
	leuc	മ	656	854	7.20E-22	-0.04	0.07	·	SUCCINYL-COA SYNTHETASE, ALPHA CHAIN: CHAIN: A; SUCCINYL-COA SYNTHETASE, BETA CHAIN; CHAIN: B:	LIGASE, GTP-SPECIFIC LIGASE, GTP-SPECIFIC
1	leyz	⋖	533	953	6.50E-30	0.51	66.0		PHOSPHORIBOSYLGLYCINA MIDE FORMYLTRANSFERASE 2; CHAIN: A, B;	TRANSFERASE TRANSFORMYLASE, PURINE BIOSYNTHESIS, ATP-GRASP
	leyz	<	534	098	1.40E-39	0.21	0.11		PHOSPHORIBOSYLGLYCINA MIDE FORMYLTRANSFERASE 2; CHAIN: A, B;	TRANSFERASE TRANSFORMYLASE, PURINE BIOSYNTHESIS, ATP-GRASP
	1gso	V	531	868	3.60E-54	0.5	0.37		GLYCINAMIDE RIBONUCLEOTIDE SYNTHETASE; CHAIN: A;	LIGASE PURD GEN PRODUCT; GAR- SYN, GLYCINAMIDE RIBONUCLEOTIDE SYNTHETASE, ATP-GRASP, 2 PURINE DE NOVO BIOSYNTHETIC PATHWAY, SUBSTRATE CHANNELING
	liow		530	859	1.80E-43	-0.07	0.18		D-ALA':D-ALA LIGASE; CHAIN: NULL;	LIGASE DD-LIGASE, DDLB; GLYCOGEN PHOSPHORYLASE, LIGASE, CELL WALL, PEPTIDOGLYCAN 2 SYNTHESIS, VANCOMYCIN, ADP BINDING
	2scu	8	656	844	1.80E-27	0.04	0.07		SUCCINYL-COA LIGASE; CHAIN: A. D. SUCCINYL- COA LIGASE; CHAIN: B, E;	LIGASE SCS; SCS; CITRIC ACID CYCLE, HETEROTETRAMER, LIGASE
	1660	V	7	443	0	1.05	_		ELONGATION FACTOR EEFIA; CHAIN: A; ELONGATION FACTOR EEFIBA; CHAIN: B;	TRANSLATION PROTEIN-PROTEIN COMPLEX
	laox	<	119	313	9.00E-28	99.0	_		INTEGRIN ALPHA 2 BETA; CHAIN: A, B;	INTEGRIN INTEGRIN, CELL ADHESION, GLYCOPROTEIN
	laox	۷.	120	316	9.00E-28			84.82	INTEGRIN ALPHA 2 BETA; CHAIN: A, B;	INTEGRIN INTEGRIN, CELL ADHESION, GLYCOPROTEIN
776	Iaox	A	321	504	7.20E-28	0.74	-		INTEGRIN ALPHA 2 BETA;	INTEGRIN INTEGRIN, CELL

PDB Chain Start LD LD AA	 1	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound CHAN: A B:	PDB annotation ADHESION, GLYCOPROTEIN
A 122 293	293	$\overline{}$	9.00E-24	97.0	_		VON WILLEBRAND FACTOR; CHAIN: A, B;	COLLAGEN-BINDING COLLAGEN- BINDING, HEMOSTASIS, DINUCLEOTIDE BINDING FOLD
A 122 305	305		9.00E-24			94.53	VON WILLEBRAND FACTOR; CHAIN: A, B;	COLLAGEN-BINDING COLLAGEN- BINDING, HEMOSTASIS, DINUCLEOTIDE BINDING FOLD
A 324 492	492	T	1.80E-21	89.0	-		VON WILLEBRAND FACTOR; CHAIN: A, B;	COLLAGEN-BINDING COLLAGEN- BINDING, HEMOSTASIS, DINUCLEOTIDE BINDING FOLD
lauq 110 317	317	T	3.60E-32	0.42	69:0		AT DOMAIN OF VON WILLEBRAND FACTOR; CHAIN: NULL;	WILLEBRAND WILLEBRAND, BLOOD COAGULATION, PLATELET, GLYCOPROTEIN
lauq 311 509	209	1	7.20E-29			86.49	AI DOMAIN OF VON WILLEBRAND FACTOR; CHAIN: NULL;	WILLEBRAND WILLEBRAND, BLOOD COAGULATION, PLATELET, GLYCOPROTEIN
lauq 313 508	208		7.20E-29	0.78	-		A1 DOMÁIN OF VON WILLEBRAND FACTOR; CHAIN: NULL;	WILLEBRAND WILLEBRAND, BLOOD COAGULATION, PLATELET, GLYCOPROTEIN
1ck4 A 122 309	309		1.80E-28	0.39	-		INTEGRIN ALPHA-1; CHAIN: A. B;	STRUCTURAL PROTEIN I-DOMAIN, METAL BINDING, COLLAGEN, ADHESION
1ck4 A 324 500	200		3.60E-30	0.85	-		INTEGRIN ALPHA-1; CHAIN: A, B;	STRUCTURAL PROTEIN I-DOMAIN, METAL BINDING, COLLAGEN, ADHESION
1fns A 119 314	314		7.20E-31	0.27	0.75		IMMUNOGLOBULIN NMC4 IGG1; CHAIN: L; IMMUNOGLOBULIN NMC4 IGG1; CHAIN: H; VON WILLEBRAND FACTOR; CHAIN: A;	IMMUNE SYSTEM VON WILLEBRAND FACTOR, GLYCOPROTEIN IBA (A:ALPHA) BINDING, 2 COMPLEX (WILLEBRAND/IMMUNOGLOBULIN), BLOOD COAGULATION TYPE 3 2B VON WILLEBRAND DISEASE
Ifns A 319 507	 507		3.60E-28	0.84	-		IMMUNOGLOBULIN NMC-4 IGG1; CHAIN: L; IMMUNOGLOBULIN NMC-4 IGG1; CHAIN: H; VON WILLEBRAND FACTOR; CHAIN: A;	IMMUNE SYSTEM VON WILLEBRAND FACTOR, GLYCOPROTEIN IBA (A:ALPHA) BINDING, 2 COMPLEX (WILLEBRAND/IMMUNOGLOBULIN), BLOOD COAGULATION TYPE 3 2B VON WILLEBRAND DISEASE
lido 124 267	267	, ,	2.60E-30	0.7	-		INTEGRIN; CHAIN: NULL;	CELL ADHESION PROTEIN A-DOMAIN INTEGRIN, CELL ADHESION PROTEIN, GLYCOPROTEIN, EXTRACELLULAR 2 MATRIX,

PDB Chain Start ID ID AA		Start		End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
										CYTOSKELETON
1ido 124 307	307	307		l	5.40E-27	0.62	_		INTEGRIN; CHAIN: NULL;	CELL ADHESION PROTEIN A-DOMAIN INTEGRIN, CELL ADHESION PROTEIN, GLYCOPROTEIN, EXTRACELLULAR 2 MATRIX, CYTOSKELETON
lido 324 498 9	498	498	 	9	9.10E-44			96.71	INTEGRIN; CHAIN: NULL;	CELL ADHESION PROTEIN A-DOMAIN INTEGRIN, CELL ADHESION PROTEIN, GLYCOPROTEIN, EXTRACELLULAR 2 MATRLX, CYTOSKELETON
1ido 325 497 9.	497	497	 	6	9.10E-44	0.57	-		INTEGRIN; CHAIN: NULL;	CELL ADHESION PROTEIN A-DOMAIN INTEGRIN, CELL ADHESION PROTEIN, GLYCOPROTEIN, EXTRACELLULAR 2 MATRIX, CYTOSKELETON
lido 326 498 9.0	498	498	ļ — — — —	0.6	9.00E-26	0.54	-		INTEGRIN; CHAIN: NULL;	CELL ADHESION PROTEIN A-DOMAIN INTEGRIN, CELL ADHESION PROTEIN, GLYCOPROTEIN, EXTRACELLULAR 2 MATRIX, CYTOSKELETON
11fa A 123 268 3.6	123 268	268	 	3.6	3.60E-23	0.42	_		CD11A: 1LFA 5 CHAIN: A. B. 1LFA 6	CELL ADHESION LFA-1, ALPHA- L'BETA-2 INTEGRIN, A-DOMAIN; ILFA 8
11fa A 323 501 3.0	323 501	201	 	m	3.60E-24			90.24	CD11A; ILFA 5 CHAIN: A, B; ILFA 6	CELL ADHESION LFA-1, ALPHA- L\BETA-2 INTEGRIN, A-DOMAIN; ILFA 8
11fa A 326 503 3.	326 503	503		3.	3.60E-24	0.85			CD11A; ILFA 5 CHAIN: A, B; ILFA 6	CELL ADHESION LFA-1, ALPHA- L'BETA-2 INTEGRIN, A-DOMAIN; ILFA 8
309	. 122 309	309		7	7.20E-28	0.7	_		ALPHAI BETAI INTEGRIN; CHAIN: A; ALPHAI BETAI INTEGRIN; CHAIN: B;	CELL ADHESION INTEGRIN, CELL ADHESION
1qc5 A 324 499 1.	324 499	499		-	1.60E-29	1.09	-		ALPHAI BETAI INTEGRIN; CHAIN: A; ALPHAI BETAI INTEGRIN; CHAIN: B;	CELL ADHESION INTEGRIN, CELL ADHESION
1b3u A i 343 3	343	 	 	<u> m</u>	3.60E-33	0.02	0.17		PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT
lee4 A 1 322 3	1 322	Н	Н	3	3.60E-49	0.52	_		KARYOPHERIN ALPHA;	TRANSPORT PROTEIN SERINE-RICH

PDB annotation	RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA, NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN, C2-DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSIS	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN, C2-DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSIS
	RNA P PROTE	TRANS RNA P PROTE	NUCLI KARY IMPOF LOCAI ARMA AUTO REGU	NUCL KARY IMPOF LOCA ARMA AUTO REGU	STRU(REPE/ STRU(ARM/ REPE/ CYTO	ARM/ REPE, CYTO	ENDO SYNA EXOC 2 REL ENDO	ENDO SYNA EXOC 2 REL ENDO
Coumpound	CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	IMPORTIN ALPHA; CHAIN: A;	IMPORTIN ALPHA; CHAIN: A;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	SYNAPTOTAGMIN I; CHAIN: A;	SYNAPTOTAGMIN I; CHAIN: A;
SeqFold score			52.47						
PMF score		0.17		1	0.98	0.94	0.75	-	_
Verify		0.03		0.55	0.26	0.3	0.21	0.41	0.59
PSI- BLAST		5.40E-10	1.40E-51	1.40E-51	3.60E-39	1.30E-29	5.40E-31	2.60E-39	3.60E-24
End		347	344	343	347	336	347	265	264
Start		233	_	8	26	_	89	140	143
Chain 1D		<	<	<				∢	∀
PDB ID		lee4	lial	lial	2bct	3bct	3bct	1byn	1byn
SEQ ID		777	777	777	777	777	777	779	779

						_						
PDB annotation	ENDOCYTOSIS/EXOCYTOSIS BETA SANDWICH, CALCIUM ION, C2 DOMAIN	ENDOCYTOSIS/EXOCYTOSIS BETA SANDWICH, CALCIUM ION, C2 DOMAIN					COMPLEX (ZINC FINGERDNA) COMPLEX (ZINC FINGERDNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	CONTRACTILE LIM DOMAIN, CRP, NMR, MUSCLE DIFFERENTIATION. CONTRACTILE	SIGNALLING PROTEIN BINDING PROTEIN, CYTOKINE, SIGNALLING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	SYNAPTOTAGMIN III; CHAIN: A;	SYNAPTOTAGMIN III; CHAIN: A;	CALCIUMPHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	CRP1; CHAIN: A;	TUMOR NECROSIS FACTOR RECEPTOR: CHAIN: A, B:	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score			130.23				70.71			58.15	\$7.16	
PMF	_			-	_			0.03	90:00			-
Verify score	0.51	0.43		0.35	0.56			-0.31	-0.2			0.32
PSI- BLAST	2.60E-73	9.00E-55	2.60E-41	2.60E-41	3.60E-24		7.20E-31	7.20E-22	3.60E-26	7.80E-13	0.00026	3.60E-50
End	376	387	266	264	264		227	91	691	277	186	197
Start AA	140	143	136	140	143		145	61	95	88	<u>e</u>	116
Chain ID	V	V					<	<	¥	V	<	U
PDB LD	nbp1	ldqv	lrsy	lisy	Irsy		lath	lalh	lalh	168t	lext	1me y
SEQ ED	977	779	779	779	779		783	783	783	783	783	783

PDB annotation		(ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	Chycrat craincrine COMPLEY	CRISIAL SINOCIORE, COMILEO (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	NTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	CINC FINGENCIANA)	COMPLEX (ZINC FINGER/DNA) ZINC ENGER PROTEIN-DNA	INTERACTION PROTEIN DESIGN 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION
Coumpound			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA CHAIN A B D E	CONSENSUS ZINC FINGER	PROTEIN: CHAIN: C. F. G.			DNA; CHAIN: A, B, D, E,	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA CHAIN A B D E	CONSENSUS ZINC FINGER	PROTEIN, CHAIN: C, F, G,			DNA, CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		6	DNA; CHAIN: A, B, D, E;	DECTRIBITION OF DECEMBERS OF THE OFFICE OF THE OFFICE OF THE OFFICE OFFI	I NOTELIN, CITAIN: C, 1, C,		TEIIIA: CHAIN: A. D. 5S	RIBOSOMAL RNA GENE,	CHAIN: B, C, E, F;	
SeqFold											92.4																							87.93			
PMF			_			-	-								0.03	3				1					0.03					0.74							
Verify			0.54			0.47	÷.								0.15	3				0.43					-0.51					0.08							
PSI- BLAST			3.60E-50			1 305 50	1.305-30				1.30E-50				3 605 30	70-700-0				1.80E-46					5.40E-42					1.10E-45				1 30F-58	2		
End	{		225			263	667				254				10	:				276		_			141					691				275	<u>;</u>	_	
Start	ŧ		144			173	7/1	•			172				10	9				200					89					94				116	:		
Chain	3		S				ر				C				,	ر				ပ					O					ပ				A	:		
PDB			l ac	_			e ;	<u>`</u>			Ime	^			1) 	`			- E	>				Ime	^				line	<u> </u>			1166	2		
SEQ	ģ	2	783			\top	- /8/ 				783	_			107	6/				783					783					783				783	3		

PDB annotation	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGILI ATION/DNA) YING-YANG I	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(IRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING BROTEIN (DNA) EIVE ENIGER CIT CIT	FROI EIN/DINA) FIVE-FINGER GEI; GEI.	ZINC FINGER, COMPLEX (DNA-	COMPLEX (DNA BINDING	PROTEIN/DNA) FIVE-FINGER GLI: GLI.	The conference of the contraction of the contractio	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-FINGER GLI; GLI,	ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRIS PS	INITIATOR ELEMENT DNA:	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;				YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;				ZINC FINGER PROTEIN GLII;	CHAIN, A, DIVA, CHAIN, C,	ä	7TNC FINGER PROTEIN GI 11:	CHAIN: A: DNA: CHAIN: C.	fo	ý	ZINC FINGER PROTEIN GLII	CHAIN: A; DNA; CHAIN: C,	D;	
SeqFold score		85.55							_																				86.72										
PMF score								0.87							1							0.46										-	•			6.0			
Verify score								0.1							-0.02							-0.12										0 14				0.23			
PSI- BLAST		3.90E-51						9.10E-47							3.90E-51							3.60E-32							3.90E-39			3 90E-59				1.30E-56			
End		226						225							254							197							722			255				267			
Start AA		116						121							149							20							91			117				145			
Chain ID		၁						ပ							၁							ပ							∢			V				A			
PDB ID		1 ubd						l ubd							pqnl							lubd		_				:	1187			2gli	0			2gli			
SEQ ID NO:		783						783	_						783					_		783						107	60/		_	783				783			

																			_				_	_	
PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	-	TRANSFERASE ATK, AMGX1, BPK, TYROSINE KINASE, X-LINKED	AGAMMAGLOBULINEMÍA, XLA, BTK, SH3 2 DOMAIN, TRANSFERASE	COMPLEX (ADAPTOR PROTEINE) ASH GROWTH	FACTOR RECEPTOR-BOUND PROTEIN	Z; CUMPLEA (ADAF FOR PROTEIN/PEPTIDE), SH3 DOMAIN, 2	GUANINE-NUCLEOTIDE RELEASING FACTOR	COMPLEX (TRANSFERASE/PEPTIDE)	COMPLEA (TRANSDUCTION, 2 SH3 DOMAIN	COMPLEX (SIGNAL	TRANSDUCTION/PEPTIDE) COMPLEX	SH3 DOMAIN	SIGNAL TRANSDUCTION ADAPTOR	SH2, SH3 1GR1 14	CIRCULAR PERMUTANT PWT;	CIRCULAR PERMUTANT, SH3 DOMAIN, CYTOSKELETON	CYTOSKELETON CYTOSKELETON,	MEMBRANE, SH3 DOMAIN	SIGNAL TRANSDUCTION PROTEIN	SKC-HUMULUGY 3 (SH3) DUMAIN, PEPTIDE-BINDING PROTEIN, ISEM 18	2 GUANINE NUCLEOTIDE EXCHANGE	FACTOR 1SEM 19		ENDOCYTOSIS/EXOCYTOSIS NSECT; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT
Coumpound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;		BRUTON'S TYROSINE KINASE; CHAIN: NULL;		GRB2; CHAIN: A; SOS;	CHOIN: B,			ABL TYROSINE KINASE,	CHAIN: A, C, E, O, FEF 11DE P41; CHAIN: B, D, F, H;	GRB2; CHAIN: A; SOS-1;	CHAIN: B;		GROWTH FACTOR BOUND	PROTEIN 2; IGRI 5 CHAIN: A, B; IGRI 6	ALPHA SPECTRIN; CHAIN:	NULL;	ALPHA II SPECTRIN; CHAIN:	A,	SEM-5: 1SEM 3 CHAIN: A, B;	ISEM 5 10-RESIDUE PROLINE-RICH PEPTIDE	FROM MSOS ISEM 8 CHAIN:	C, D ISEM 10		SYNTAXIN BINDING PROTEIN I; CHAIN: A; SYNTAXIN IA; CHAIN: B;
SeqFold score																									
PMF score	0.1		8.0		0.83				0.25		0.87			0.12		0.59		0.71		0.59					0.01
Verify score	-0.43		0.42		0.35				0.08		9.0			0.12		0.2		-0.25		0.21					-0.67
PSI- BLAST	3.60E-26	ļ	1.00E-09		2.60E-11				3.90E-10		9.10E-12			1.80E-09		2.60E-11		1.30E-10		5.40E-11					1.00E-05
End	140		403		402				403		403			400		403		403		400					281
Start	26		340		348				350		348			343		344		349		349					160
Chain 19	<				A				A		A			٧				4		¥					<u> </u>
PDB TD	2gli		Na v		laze				1bbz		1gbq			lgi.)	Ipwt		1qk	3	Sem					1dn1
SEQ ID	783		784		784				784		784			784		784		784		784		_			785

PDB annotation	HYDROLASE HOMODIMER, ALPHA/BETA HYDROLASE FOLD, DISUBSTITUTED UREA 2 INHIBITOR	HYDROLASE LIPASE		HYDROLASE PSEUDOMONADACEAE, CIS-PEPTIDE, CLOSED CONFORMATION, 2 HYDROLASE, LID	HYDROLASE ALPHA BETA HYDROLASE FOLD, PROLINE, PROLYL AMINOPEPTIDASE, 2 SERRATIA, IMINOPEPTIDASE	LIPASE LIPASE, LIPASE, HYDROLASE, PSEUDOMONADACEAE, COVALENT INTERMEDIATE, 2 TRIGLYCERIDE ANALOGUE. ENANTIOSELECTIVITY	SIGNAL TRANSDUCTION SIGNAL TRANSDUCTION, SOS, PLECKSTRIN HOMOLOGY (PH) DOMAIN	SIGNAL TRANSDUCTION PROTEIN	TRANSPORT PROTEIN RHO-GTPASE EXCHANGE FACTOR, TRANSPORT PROTEIN	TRANSPORT PROTEIN RHO-GTPASE EXCHANGE FACTOR, TRANSPORT PROTEIN	GENE REGULATION SON OF SEVENLESS PROTEIN; GUANINE NUCLEOTIDE EXCHANGE FACTOR, GENE REGULATION
Coumpound	EPOXIDE HYDROLASE; CHAIN: A, B;	LIPASE, GASTRIC; CHAIN: A, B;	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED ILPB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) ILPB 4	TRIACYLGLYCEROL HYDROLASE; CHAIN: D; TRIACYLGLYCEROL HYDROLASE; CHAIN: E;	PROLYL AMINOPEPTIDASE; CHAIN: A;	TRIACYL-GLYCEROL- HYDROLASE: CHAIN: D, E;	SOS1; CHAIN: NULL;	BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	PIX; CHAIN: A;	PIX; CHAIN: A;	HUMAN SOS 1; CHAIN: A;
SeqFold score											
PMF score	0.13	0	0	-0.03	0	0.01	0.49	-	0.89	9.0	96.0
Verify score	-0.11	-0.32	-0.29	0.1	-0.16	-0.09	0.32	0.39	-0.25	-0.46	-0.1
PSI- BLAST	1.10E-37	1.60E-07	1.40E-06	3.60E-11	1.60E-28	1.60E-11	1.20E-23	9.00E-09	2.60E-41	7.20E-23	1.80E-16
End	258	121	149	203	242	191	576	819	463	462	570
Start	14	5	30	25	∞	25	462	728	261	267	261
Chain ID	В	Y	B	Ω	<	Q			<	<	4
PDB ID	leki	1hlg	11pb	Iqge	lqtr	4lip	lawe	1btn	1by1	1by1	1dbh
SEQ ID	786	786	786	786	786	786	788	788	788	788	788

	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	8	8	۷ ۲	V V	BLAST	score	score	score		
	1dbh	 	263	576	7.80E-58	0.12	_		HUMAN SOS 1; CHAIN: A;	GENE REGULATION SON OF SEVENLESS PROTEIN; GUANINE NUCLEOTIDE EXCHANGE FACTOR, GENE REGULATION
T^{-}	Idro		736	820	9.10E-09	0.22	0.21		BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7	CYTOSKELETON
	ldvp	4	618	674	9.00E-11	0.07			HEPATOCYTE GROWTH FACTOR-REGULATED TYROSINE CHAIN: A;	TRANSFERASE HRS, HRS, VHS, FYVE, ZINC FINGER, SUPERHELIX
T	1f5x	<	260	454	5.40E-24	0.14			RHO-GEF VAV, CHAIN: A;	SIGNALING PROTEIN 11 ALPHA- HELICES
	lfao	⋖	726	816	3.60E-11	0.52	0.94		DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN
	1fb8	∀	484	574	1.00E-09	-0.4	0.29		DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	SIGNALING PROTEIN DAPPI, PHISH, BAM32; PLECKSTRIN. 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN
788	11b8	⋖	719	817	2.60E-18	0.63	66.0		DUÁL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	SIGNALING PROTEIN DAPP I, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN
788	1fb8	<	727	816	1.80E-11	0.67	86.0		DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	SIGNALING PROTEIN DAPP1, PHISH, BAM32, PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN
	1fgy	٧	484	574	9.10E-08	-0.18	0.21		GRP1; CHAIN: A;	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN
	1fgy	∢	725	821	1.10E-15	0.27	0.98		GRP1; CHAIN: A;	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN

PDB annotation	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN		SIGNAL TRANSDUCTION IRS-1; BETA-SANDWHICH, SIGNAL TRANSDUCTION	TRANSPORT PROTEIN FYVE DÖMAIN, ENDOSOME MATURATION, INTRACELLULAR TRAFFICKING, 2 TRANSPORT PROTEIN	COMPLEX (GTP-BINDING/EFFECTOR) RAS-RELATED PROTEIN RAB3A; COMPLEX (GTP-BINDING/EFFECTOR), G PROTEIN, EFFECTOR, RABCDK, 2 SYNAPTIC EXOCYTOSIS, RAB PROTEIN, RAB3A, RABPHILIN	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKINES, THIRD 2 N-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKINES, THIRD 2 N-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKINES, THIRD 2 N-TERMINAL DOMAIN, TRANSMEMBRANE, GLYCOPROTEIN
Coumpound	GRP1; CHAIN: A:	PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS IPLS 4 (INS(G105-LEHHHHHH)) (NMR, 25 STRUCTURES) IPLS 5	INSULIN RECEPTOR SUBSTRATE 1; CHAIN: A, B;	PHOSPHATIDYLINOSITOL-3- PHOSPHATE BINDING FYVE CHAIN: A;	RAB-3A; CHAIN: A; RABPHILIN-3A; CHAIN: B;	GP130; CHAIN: NULL;	GP130; CHAIN: NULL;	GP130; CHAIN: NULL;
SeqFold								
PMF	90.0	0.93	0.76	0.94	0.1	0.09	0.13	-0.01
Verify score	-0.15	0.1	0.64	0.04	0.11	-0.08	-0.09	0.14
PSI- BLAST	7.80E-16	1.80E-11	3.60E-05	7.20E-08	3.90E-21	1.10E-08	3.60E-11	7.20E-11
End AA	814	819	918	671	674	360	477	580
Start	729	728	725	620	290	289	387	484
Chain ID	∢		A	٧	В			
PDB ID	Ifgy	1 pis	Iqqg	lvfy	1zbd	1bj8	16j8	15j8
SEQ ID NO:	788	788	788	788	788	789	789	789

PDB annotation	CONNECTIN, FIBRONECTIN, TITIN, CONNECTIN, FIBRONECTIN TYPE III	CONNECTIN, FIBRONECTIN, TITIN, CONNECTIN, FIBRONECTIN TYPE III	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN		HORMONE/GROWTH FACTOR/HORMONE RECEPTOR 4- HELICAL BUNDLE, ALPHA HELICAL BUNDLE, TERNARY COMPLEX, FN 2 III DOMAINS, BETA SHEET DOMAINS, CYTOKINE-RECEPTOR COMPLEX			
Coumpound	TITIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	GP130; CHAIN: A, B;	GP130; CHAIN: A, B;	GP130; CHAIN: A, B;	NEURAL ADHESION MOLECULE DROSOPHILA NEUROGLIAN (CHYMOTRYPTIC FRAGMENT CONTAINING THE ICFB 3 TWO AMINO PROXIMAL FIBRONECTIN TYPE III REPEATS ICFB 4 (RESIDUES 610 - 814)) ICFB 5	PLACENTAL LACTOGEN; CHAIN: A; PROLACTIN RECEPTOR; CHAIN: B, C;	CELL ADHESION PROTEIN FIBRONECTIN CELL- ADHESION MODULE TYPE III-10 1FNA 3	CELL ADHESION PROTEIN FIBRONECTIN CELL- ADHESION MODULE TYPE III-10 1FNA 3	CELL ADHESION PROTEIN FIBRONECTIN CELL-
SeqFold score										
PMF	0.11	0.25	90.0	-0.14	-0.03	0.24	61.0	0.19	0.22	0.41
Verify score	-0.31	-0.02	-0.19	0.09	0.11	0.03	-0.26	-0.14	0.04	0.07
PSI- BLAST	3.60E-07	3.60E-10	7.20E-21	9.00E-15	1.80E-18	1.80E-26	1.60E-14	3.60E-07	1.10E-11	1.30E-12
End	360	878	360	490	595	583	359	360	582	286
Start	288	485	195	288	386	385	196	295	492	504
Chain ID			<	¥	4		a			
PDB ID	lbpv	1bpv	1bqu	1bqu	1bqu	1cfb	1166	lfna	1 fna	1 fna
SEQ ID	789	789	789	789	789	789	789	789	789	189

SEQ	PDB	Chain	Start	End	-ISd	Verify	PMF	SeqFold	Coumpound	PDB annotation
ΘÖ	<u>0</u>	<u>e</u>	VΥ	¥	BLAST	score	score	score		
									ADHESION MODULE TYPE III-10 1FNA 3	
789	lfnf		192	582	5.40E-39	-0.02	0.51		FIBRONECTIN; 1FNF 6 CHAIN: NULL; 1FNF 7	CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX 1FNF 18
789] If		194	588	5.40E-39			120.84	FIBRONECTIN; 1FNF 6 CHAIN: NULL: 1FNF 7	CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX IFNF 18
789	Ifuf		288	674	9.00E-38	-0.03	0.31		FIBRONECTIN; 1FNF 6	CELL ADHESION PROTEIN RGD,
789	1fnf		386	752	9.00E-39	0.02	-0.07		FIBRONECTIN; IFNF 6	CELL ADHESION PROTEIN RGD
789	1 finh	<	961	473	1.10E-19	0.11	0.07		FIBRONECTIN; CHAIN: A;	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING
789	1fnh	4	389	673	7.20E-32	0.05	0.55		FIBRONECTIN; CHAIN: A;	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING
789	Har	٧	893	1462	0	99.0	_		LAR; CHAIN: A, B;	HYDROLASE TYROSINE PHOSPHATEASE, LAR PROTEIN
789	llar	В	782	1164	3.60E-81	0.34	_		LAR; CHAIN: A, B;	HYDROLASE TYROSINE PHOSPHATEASE, LAR PROTEIN
789	Har	В	924	1462	0	0.72	_		LAR; CHAIN: A, B;	HYDROLASE TYROSINE PHOSPHATEASE, LAR PROTEIN
789	1mfn		292	474	5.40E-23	0.12	0.16		FIBRONECTIN, CHAIN: NULL;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HFPARIN, RIDING, GI, YCOPROTEIN
789	lmfn		388	582	1.80E-25	-0.13	60.0		FIBRONECTIN; CHAIN: NULL;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN
789	1923	×.	290	478	1.10E-17	0.18	0.1		INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN
789	1983	⋖	388	588	1.40E-25	0.28	-1.41		INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN
789	1983	V	390	584	2.60E-29	0.3	0.92		INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN
789	1qr4	Y	292	480	9.00E-17	0.12	0.18		TENASCIN; CHAIN: A, B;	STRUCTURAL PROTEIN TENASCIN,

PDB annotation	FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TENASCIN, FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN	RECEPTOR DI; RECEPTOR, PHOSPHATASE, SIGNAL TRANSDUCTION, ADHESION, 2 HYDROLASE	RECEPTOR DI; RECEPTOR, PHOSPHATASE, SIGNAL TRANSDUCTION, ADHESION, 2 HYDROLASE				IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	TYROSINE PHOSPHATASE SYP, SHPTP-2; TYROSINE PHOSPHATASE, INSULIN SIGNALING, SH2 PROTEIN
Coumpound		TENASCIN; CHAIN: A, B;	RECEPTOR PROTEIN TYROSINE PHOSPHATASE MU; CHAIN: A, B;	RECEPTOR PROTEIN TYROSINE PHOSPHATASE MU; CHAIN: A, B;	CELL ADHESION PROTEIN TENASCIN (THIRD FIBRONECTIN TYPE III REPEAT) ITEN 3	CELL ADHESION PROTEIN TENASCIN (THIRD FIBRONECTIN TYPE III REPEAT) ITEN 3	GLYCOPROTEIN FIBRONECTIN (TENTH TYPE III MODULE) (NMR, 36 STRUCTURES) 1TTF 3	FC GAMMA RIIB; CHAIN: A;	FIBRONECTIN; CHAIN: A;	FIBRONECTIN, CHAIN: A;	SHP-2; CHAIN: A, B;
SeqFold score			405.81								
PMF		0.41			0.03	0.03	0.31	-0.2	0.98	0.16	_
Verify score		0.32		0.85	0.08	0.08	0.16	0.03	0.35	0.31	74.0
PSI- BLAST		1.305-20	1.40E-89	1.40E-89	1.60E-07	1.30E-08	5.40E-13	1.40E-10	1.60E-05	1.80E-08	1.40E-72
End		584	1166	1165	584	584	582	163	480	584	1164
Start		390	688	891	487	502	485	27	391	490	835
Chain ID		<	∀	A				٧	<	V	¥
PDB ID		lqr4	Irpm	Irpm	Iten	Iten	lπf	2fcb	2fnb	2fnb	2shp
SEQ ID	Ž	789	682	789	789	789	789	789	789	789	789

PDB Chain Start End I	Start End AA AA	End		_ <u>~</u>	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
1b8q A 48 176 1.40E-11	48 176	176		1.40E-11				58.72	NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE: CHAIN: B:	OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE
1b8q A 52 171 1.40E-11 (52 171 1.40E-11	171 1.40E-11	1.40E-11			0.46	0.23		NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE
1be9 A 45 163 9.00E-16	45 163	163		9.00E-16	1			57.29	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION
lbc9 A 49 143 9.00E-16 1	49 143 9.00E-16	143 9.00E-16	9.00E-16			1.06	-		PSD-95; CHAIN: A; CRIPT; CHAIN: B;	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION
1btn 299 388 3.60E-10 -c	388 3.60E-10	388 3.60E-10	3.60E-10		۲	-0.05	0.04		BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	SIGNAL TRANSDUCTION PROTEIN
Ifao A 295 385 5.40E-09 0.24	295 385 5.40E-09	385 5.40E-09	5.40E-09		0.7	4	-0.13		DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	SIGNALING PROTEIN DAPP1, PHISH, BAM32; PLECKSTRIN, 3-PHOSPHOINOSITIDES, INOSITOL TETRAKISPHOSPHATE 2 SIGNAL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN
1fgy A 199 267 0.0052 -0.65	199 267 0.0052	267 0.0052	0.0052		9.0-	δ.	0.05		GRP1; CHAIN: A;	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN
1i16 27 138 3.60E-11 0.09	138 3.60E-11	138 3.60E-11	3.60E-11		0.0	0	0.43		INTERLEUKIN 16; CHAIN: NULL;	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN
1qau A 54 166 3.60E-10 0.5	54 166 3.60E-10	166 3.60E-10	3.60E-10		0.5		66:0		NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130); CHAIN: A;	OXIDOREDUCTASE BETA-FINGER
lqau A 56 168 2.60E-23 0.47	56 168 2.60E-23	168 2.60E-23	2.60E-23		0.4	17	0.94		NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130); CHAIN: A;	OXIDOREDUCTASE BETA-FINGER
lqav A 54 140 3.90E-23	54 140	140		3.90E-23				77.08	ALPHA-I SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES I-130); CHAIN: B;	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER
1qav A 56 136 3.90E-23 0.	56 136 3.90E-23	136 3.90E-23	3.90E-23		o l	0.87			ALPHA-I SYNTRÖPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER

PDB annotation		MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER	RNA-BINDING PROTEINIRNA TRA PRE-MRNA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX	GENE REGULATION/RNA POLY(A)	BINDING PROTEIN 1, PABP 1; RRM,	PROTEIN-RNA COMPLEX, GENE	KEGULATION/KNA		RNA BINDING PROTEIN RNA- BINDING DOMAIN	NUCLEAR PROTEIN	HETEROGENEOUS NUCLEAR	RIBONUCLEOPROTEIN AI, NUCLEAR	PROTEIN, HNRNP, RBD, RRM, RNP,	RNA BINDING, 2	KIBUNUCLEURUIEIIN	NUCLEAR PROTEIN HETTEROGENEOUS NUCLEAR	RIBONUCLEOPROTEIN A1, NUCLEAR	PROTEIN, HNRNP, RBD, RRM, RNP,	RNA BINDING, 2 RIBONLICI EOPROTEIN	COMPLEX	(RIBONUCLEOPROTEIN/RNA)					RNA BINDING PROTEIN RNA- BINDING DOMAIN
Coumpound	(RESIDUES 1-130); CHAIN: B;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B;	SXL-LETHAL PROTEIN; CHAIN: A, B; RNA (5'- R(P*GP*UP*UP*UP*U P*UP*UP*UP*UP*U)- CHAIN: P, Q;	POLYDENYLATE BINDING	PROTEIN I; CHAIN: A, B, C,	D, E, F, G, H; RNA (5'-	R(*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN:	M, N, O, P, Q, R, S, T,	HU ANTIGEN C; CHAIN: A;	HNRNP AL CHAIN: NULL;						HNRNP AI; CHAIN: NULL;				UIA SPLICEOSOMAL	PROTEIN; IURN 5 CHAIN: A,	B, C; IURN 6 RNA 21MER	HAIRPIN (5'-	(AP*AP*UP*CP*CP*AP*UP*U	P* IURN II CHAIN: P, Q, R	MUSASHII: CHAIN: A;
SeqFold score						_																				
PMF score		-	0.81	66.0					0.71	0.63						0.99				0.94						0.88
Verify score		0.93	0.84	0.41					0.39	0.48						89.0				0.55				<u>.</u>		0.82
PSJ- BLAST		3.60E-16	1.80E-16	5.40E-18					1.30E-15	9 00E-25						1.80E-15				2.60E-15						1.60E-15
End		139	108	100	<u>}</u>				105	103					_	107				112	:					103
Start		95	33	~	,				34	_	•					34				33	}					35
Chain ID		¥	∀	A	:				4											4	:					<
PDB ID		Iqav	167f	ivi	· ·				148z	1 ha 1						Ihal				- III						2mss
SEQ ID	2	793	794	707	<u> </u>				794	704			_			194				794	· -	_				794

PDB annotation	RNA-BINDING DOMAIN RNA- BINDING DOMAIN, ALTERNATIVE SPLICING	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	RNA BINDING DOMAIN RNA BINDING DOMAIN, RBD, RNA RECOGNITION MOTIF, RRM, 2 SPLICING INHIBITOR, TRANSLATIONAL INHIBITOR, SEX 3 DETERMINATION, X CHROMOSOME DOSAGE COMPENSATION	COMPLEX (TRANSCRIPTION REGULATION/DNA) SREBP-1A; STEROL REGULATORY ELEMENT BINDING PROTEIN, 2 BASIC-HELIX- LOOP-HELIX-LEUCINE ZIPPER, SREBP, TRANSCRIPTION 3 FACTOR, COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) SREBP-1A; STEROL REGULATORY ELEMENT BINDING PROTEIN, 2 BASIC-HELIX- LOOP-HELIX-LEUCINE ZIPPER, SREBP, TRANSCRIPTION 3 FACTOR, COMPLEX (TRANSCRIPTION	COMPLEX (DNA-BINDING PROTEIN/DNA) MYN PROTEIN; MAX, DNA BINDING, BASIC-HELIX-LOOP- HELIX-LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) UPSTREAM
Coumpound	SEX-LETHAL PROTEIN; CHAIN: NULL;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	SEX-LETHAL; CHAIN: A, B, C,	STEROL REGULATORY ELEMENT BINDING PROTEIN IA; CHAIN: A, B, C, D: DNA; CHAIN: E, F, G, H;	STEROL REGULATORY ELEMENT BINDING PROTEIN IA; CHAIN: A, B, C, D; DNA; CHAIN: E, F, G, H;	MAX PROTEIN; CHAIN: A, C; DNA; CHAIN: B, D;	USF; CHAIN: A, B; DNA; CHAIN: C, D;
SeqFold score							
PMF	99.0	0.23	0.96	0.04	0.27	-0.03	0.07
Verify	0.31	0.68	0.61	0.23	0.14	0.35	-0.08
PSI- BLAST	1.80E-16	5.40E-30	5.40E-16	2.60E-13	1.30E-15	9.00E-15	1.80E-13
End AA	108	11	105	131	119	119	115
Start AA	33		33	59	55	57	55
Chain ID		<	<	V	B	∢	4
PDB ID	2sxl	2up1	3sxl	lam 9	laın 9	lan2	lan4
SEQ ID NO:	794	794	794	795	795	795	795

PDB annotation	STIMULATORY FACTOR 1; USF, DNA BINDING, BASIC-HELIX-LOOP-HELIX, LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)		COMPLEX (TRANSCRIPTION FACTOR MAX/DNA) TRANSCRIPTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAX/DNA)					COMPLEX (ZINC FINGER/DNA) ZINC
Coumpound		TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5'- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5'- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	TRANSCRIPTION ACTIVATION/DNA MYOD BASIC-HELIX-LOOP-HELIX (BHLH) DOMAIN IMDY 3 (RESIDUES 102 - 166) MUTANT WITH CYS 135 REPLACED BY SER IMDY 4 (C135S) COMPLEXED WITH DNA IMDY 5 (5- D(*TP*GP*AP*AP*GP*AP*GP *CP*TP*GP*TP*TP*GP*A)-3')	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C,	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E,
SeqFold score								
PMF score		0.17	0.05	11.0-	-0.19	0.55	0.42	89.0
Verify score		90:00	-0.17	0.15	0.07	0.24	0.31	0.14
PSI- BLAST		3.60E-15	7.20E-15	1.40E-14	1.40E-22	3.60E-26	5.20E-32	5.40E-44
End		119	119	411	405	433	434	433
Start AA		53	55	23	327	353	357	352
Chain ID		4	æ	æ	∢	∢	<	S
PDB ID		1hlo	1hlo	y y	lalh	lath	lalh	Ime
SEQ ID	ÖZ	795	795	795	800	800	800	800

PDB annotation				ER 3;					, E; COMPLEX (ZINC FINGER/DNA) ZINC NGER FINGER, PROTEIN-DNA
Coumpound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA, CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA, CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA, CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER
SeqFold score									
PMF score		_		_	1	1	-	-	-
Verify score		0.49	0.45	0.52	0.52	0.48	0.45	0.15	61.0
PSI- BLAST		1.80E-46	3.90E-48	2.60E-48	5.40E-47	9.00E-48	1.80E-48	9.00E-50	1.80E-50
End		461	461	489	489	517	545	573	109
Start AA		380	381	408	408	436	464	492	520
Chain ID		C	ပ	O	U	၁	U	ن د	၁
PDB ID	٨	y me	Ime y	, me	ر ب سر	Ime y	y y	y y	1mc y
SEQ B		008	800	800	800	800	008	800	800

PDB annotation	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA;	TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC	FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), MAYA POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	FULT MERASE III, 2 I RANSCAIL LION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	REGIII ATION/DNA). RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TING-TANG 1,	INITIATOR FLEMENT YYL ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(IKANSCKIPTION KEGOLATION/DINA)	COMPLEA (IRANSCALFIION
Coumpound		TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;			TFIIIA; CHAIN: A, D; 5S	CHAIN: B, C, E, F;			TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN A B.	Credit: A, B,		Olding.	YYI; CHAIN: C; ADENO-
SeqFold score													111.83																
PMF		-0.2			0.54				1									_					0.43					,	0.96
Verify score		0			0.2				0.31									-0.13					-0.16						0.15
PSI- BLAST		3.60E-11			1.80E-35				9.00E-38				7.20E-38					7.20E-38					1.30E-28						5.40E-31
End		377			498				999				741					754					433			_			461
Start AA		292			353				521				576					509					332						355
Chain ID		V			A				A				A					4					O						ပ
PDB ID		11.0			1166				9311				116					1116			_		lubd						Iubd
SEQ ID	Ž	800			800			_	800				800					800					808						80

PDB annotation	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1, CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold						
PMF score		-	-	_	0.96	0.93
Verify score		0.05	0.51	0.1	0.12	0.33
PSI- BLAST		1.30E-45	1.30E-57	7.80E-55	1.30E-53	9.00E-35
End		461	489	545	601	629
Start		357	385	434	490	528
Chain ID		S	O	O	ပ	၁
PDB ED		lubd	Iubd	lubd	lubd	lubd
SEQ ID	Ž	008	800	800	800	800

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA).	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA).	COMPLEX (TRANSCRIPTION REGULATION/DNA) YNG-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA: CHAIN: A, B:	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; ÇHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score		91.29				
PMF score			-	0.99		
Verify score	0.11		0.22	0.25	0.13	0.15
PSI- BLAST	1.30E-58	5.20E-60	3.60E-35	5.20E-60	1.60E-34	3.90E-61
End	657	658	657	713	713	741
Start AA	546	548	556	602	612	630
Chain ID	U	O	U	U	U	U
PDB ID	lubd	pqnl	lubd	lubd	pqn	1ubd
SEQ D	800	800	800	800	800	800

PDB Chain Start End PSI- Verify ID AA AA BLAST score	Start End PSI- AA AA BLAST	End PSI- AA BLAST	PSI- BLAST		Verify score	-	PMF	SeqFold	Coumpound	PDB annotation (TRANSCRIPTION REGULATION/DNA)
2gli A 367 463 1.30E-43 0.44	367 463 1.30E-43	463 1.30E-43	1.30E-43		0.44		0.93		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 380 519 3.90E-71	380 519	519		3.90E-71				105.23	ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 381 547 2.60E-70 0.18	381 547 2.60E-70	547 2.60E-70	2.60E-70		0.18		0.88		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 408 603 3.90E-71 0.06	408 603 3.90E-71	603 3.90E-71	3.90E-71		90.0		0.81		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 464 631 5.20E-73 0.15	464 631 5.20E-73	631 5.20E-73	5.20E-73		0.15		-		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 528 656 1.30E-34 0.16	528 656 1.30E-34	656 1.30E-34	1.30E-34		0.16		6.0		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 576 743 5.20E-79 0.11	576 743 5.20E-79	743 5.20E-79	5.20E-79		0.11		-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 584 712 5.40E-34 0.32	584 712 5.40E-34	712 5.40E-34	5.40E-34		0.32		-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 612 743 1.10E-33 0.41	612 743 1.10E-33	743 1.10E-33	1.10E-33		0.41		0.98		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
lcun A 91 302 0.0052 -0.2	91 302 0.0052	302 0.0052	0.0052		-0.2		0.37		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 22

Chain ID		Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
4		54	257	3.90E-59 6.50E-70	0.41	_ -		ALPHA-CATENIN, CHAIN: A, ALPHA-CATENIN, CHAIN: A; BETA-CATENIN, CHAIN: B;	BUNDLE CELL ADHESION FOUR-HELIX BUNDLE
<		367	441	0.0072	-0.65	90.0		RIBOSOME RECYCLING FACTOR; CHAIN: A;	RIBOSOME TRANSLATION, RIBOSOME, HINGE VARIABILITY
4		2	103	1.30E-06	-0.34	10:0		PREFOLDIN; CHAIN: A; PREFOLDIN; CHAIN: B; PREFOLDIN; CHAIN: C;	CHAPERONE ARCHAEAL PROTEIN
	1	3949	3984	6.50E-12	0.39	0.58		EPIDERMAL GROWTH FACTOR, CHAIN: NULL;	GROWTH FACTOR [ABU6, 20] MEGF4- 48, GROWTH FACTOR, MURINE EPIDERMAL GROWTH FACTOR, DISULFIDE 2 CONNECTIVITIES, EGF- LIKE DOMAIN, REPEAT
٠ <u>-</u>	i	3943	4033	6.50E-19	0.55	-0.12		ACTIVATED PROTEIN C; CHAIN: C, L, D-PHE-PRO- MAI, CHAIN: P;	COMPLEX (BLOOD COAGULATION/INHIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/INHIBITOR)
	1	3949	3987	1.30E-11	96.0	0.99		FACTOR VII; CHAIN: NULL;	BLOOD COAGULATION BLOOD COAGULATION, EGF, HYDROLASE, SERINE PROTEASE
₹	1	3771	3928	1.30E-26	0.26	0.4		NEUREXIN-I BETA; CHAIN: A, B, C. D. E. F. G, H;	MEMBRANE PROTEIN LECTIN-LIKE, NEUROBIOLOGY, CELL-CELL ADHESION, CELL-CELL 2 RECOGNITION, ALTERNATIVE SPLICING, MEMBRANE PROTEIN
<		3773	3934	2.60E-27	0.46	0.72		SEX HORMONE-BINDING GLOBULIN; CHAIN: A;	TRANSPORT PROTEIN SHBG; STEROID TRANSPORT, LAMININ G- LIKE DOMAIN, JELL YROLL, 2 ANDROGEN BINDING PROTEIN (ABP), SEX STEROID BINDING PROTEIN 3 (SBP)
J	1	3944	4033	3.90E-21	0.48	0.05		DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX

SEQ S D S	PDB ID	Chain D	Start	End	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
									(LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	
808	ldva	-1	4263	4344	1.30E-13	0.25	-0.18		DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN:	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX
808	ldx5		4232	4336	1.30E-11	90.0	-0.19		THROMBIN LIGHT CHAIN; CHAIN: A, B, C, D; THROMBIN HEAVY CHAIN; CHAIN: M, N, O, P; THROMBOMODULIN; CHAIN: I, J, K, L; THROMBIN INHIBITOR L-GLU-L-GLY-L- ARM; CHAIN: E, F, G, H;	SERINE PROTEINASE COAGULATION FACTOR II; ETOMODULIN, TM, CD141 ANTIGEN, EGR-CMK SERINE PROTEINASE, EGF-LIKE DOMAINS, ANTICOAGULANT COMPLEX, 2 ANTICOAGULANT COMPLEX, 2
808	ledh	<	9901	1234	3.60E-33	0.26	-		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	ledh	<	1171	1338	1.30E-32	0.22	1202.08		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	1edh	<	1279	1440	3.60E-21	0.45	6.0		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	ledh	V	1352	1547	1.60E-49	0.27	0.82		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	1edh	∢	1460	1652	1.30E-32	0.1	-1.41		E-CADHERIN; CHAIN: A. B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I

PDB annotation	AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	, B; CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN		, B: CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2. ECAD12; CADHERIN. CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	, B; CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN		L. B; CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	
Coumpound		E-CADHERIN; CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B.	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;
SeqFold score									
PMF		0.96	0.99	_	0	0.96	0.51	0.78	
Verify score		0.21	0.19	0.32	0	0.26	0.3	0.24	0.18
PSI- BLAST		3.60E-33	1.60E-28	3.60E-26	1.60E-20	3.60E-29	1.10E-50	1.30E-28	3.60E-48
End		1750	1860	1960	354	2062	2163	2264	2371
Start		1589	1690	1800	182	1898	1975	2104	2178
Chain ID		4	<	4	⋖	≺	≺	∢	<
PDB ID		ledh	ledh	ledh	ledh	ledh	ledh	ledh	1cdh
SEQ ID NO:		808	808	808	808	808	808	808	808

PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2. ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;						
SeqFold score						120.62			
PMF		_	0.86	0.93	96.0		_		0.94
Verify		0.32	0.07	0	0.1		0.47	0.27	0.31
PSI- BLAST		1.60E-35	1.80E-29	1.80E-38	5.40E-32	1.80E-57	1.80E-57	7.20E-35	5.40E-29
End		2473	2577	2683	2789	2895	2898	3003	3105
Start		2306	2414	2488	2619	2692	2693	2831	2941
Chain ID		A	<	<	<	<	∢	V	¥
PDB ID		1edh	ledh	1edh	ledh	ledh	ledh	ledh	1edh
SEQ ID	2	808	808	808	808	808	808	808	808

PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHEKIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	CELL A DHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12, CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS	AND 2, ECADI2; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS	AND 2, ECADIZ, CADHEKIN, CELL	ADHESION PROTEIN, CALCIUM	SELECTION OF SELECTION DE CATERIO	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12, CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Coumpound		E-CADHERIN; CHAIN: A, B;			E-CADHERIN; CHAIN: A, B;				E-CADHERIN: CHAIN: A. B;				C CADITERNI: CHARL A B.	E-CADRENIN; CRAIN. A, B,				E-CADHERIN; CHAIN: A, B;					E-CADHERIN; CHAIN: A, B;		_		COADURABLE CITAIN A D.	E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;	_		
SeqFold score																													_						
PMF		0.53			0.64				-141				5	0.98				-					0.3					0.83		_		0.82			
Verify score		0.38			0.16				0.57				,	C.40				99.0					0.12				000	0.29				0.17			
PSI- BLAST		7.20E-25			5.40E-28				1 80F-32				07 200 .	1.80E-48				3.60E-30					1.80E-51					1.10E-29				3.60E-29		• • •	
End		450			3197				3313	2			3,7,5	3418				3523					248					256				799			
Start		296			3046				3120	<u>.</u>			1000	2775				3355					39					406				464			
Chain TO		Y		-	4				A					∢				A					4				-	<				4			
PDB ID		1cdh			ledh				4									ledh					-Gg-		_			ledh				ledh			
SEQ ID		808			808		_		808					808	_			808					808					808				808		_	

SEQ	PDB	Chain	Start	End	PSI.	Verify	PMF	SeqFold	Coumpound	PDB annotation
e ë	9	e -	¥ ¥	ΥΥ	BLAST	score	score	score		
808	1edh	<	591	812	5.40E-22	-0.05	0.27		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN. CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	ledh	<	718	917	1.10E-55	0.33	-		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	ledh	4	854	1022	1.10E-32	0.31	-		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	ledh	∢	656	1129	9.00E-32	0.11	69'0		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	u u		3946	4022	1.80E-16	0.71	66.0		FIBRILLIN; CHAIN: NULL;	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-I FRAGMENT, MATRIX PROTEIN
808	l fsb		3949	3988	1.30E-11	1.03	0.88		P-SELECTIN; CHAIN: NULL;	CELL ADHESION PROTEIN EGF-LIKE DOMAIN, CELL ADHESION PROTEIN, TRANSMEMBRANE, 2 GLYCOPROTEIN
808	1klo		3924	4050	1.10E-13	-0.1	0.27		LAMININ; CHAIN: NULL;	GLYCOPROTEIN GLYCOPROTEIN
808	iklo Klo		3954 4201	4075	5.40E-21 3.60E-17	0.07	-0.03		LAMININ; CHAIN: NULL;	GLYCOPROTEIN GLYCOPROTEIN GLYCOPROTEIN GLYCOPROTEIN
808	Incg		1062	1127	9.00E-06	0.12	0.29		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	lncg		1167	1232	0.00014	0.13	69:0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	lncg		1350	1439	1.40E-14	0.01	0.29		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		1480	1546	0.00018	0.46	0.1		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
										INCG 13
808	Incg		155	229	3.60E-05	0.15	60:0		N-CADHERIN, INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		1599	1650	9.00E-06	-0.13	60.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		1667	1748	3.60E-06	0.35	0.31		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	lncg		1970	2061	3.60E-17	0.22	0.1		N-CADHERIN, INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	lncg		2079	2161	0.00036	0.28	0.11		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		2178	2263	3.60E-12	-0.08	0.1		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	lncg		2304	2370	1.80E-06	0.41	0.74		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		2411	2458	0.00036	-0.27	0.43		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		2593	2681	1.60E-05	0.34	0.57		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		2692	2788	1.80E-19	0.44	0.7		N-CADHERIN, INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		2913	2988	0.00054	0.35	0.39		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	lncg		3039	3106	3.60E-06	0.45	0.51		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		3120	3191	1.80E-05	90.0	0.34		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		3225	3311	1.60E-11	0.33	0.36		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	Incg		716	811	7.20E-20	0.16	8.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	1ncg		852	006	0.00018	-0.19	0.53		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	1ncg		932	1003	5.40E-05	0.38	0.64		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
808	1nci	8	5901	1129	1.80E-06	90.0	0.78		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	B	1172	1234	5.40E-05	90.0	96.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1350	1440	1.30E-13	0.43	0.22		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13

SEO	PDB	Chain	Start	End	PSI-	Verify	PMF	SeaFold	Coumpound	PDB annotation
ΘŞ	a	2	¥ —	¥ ¥	BLAST	score	score	score		
808	Inci	B	1491	1547	0.00018	-0.08	9.65		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	83	1667	1750	9.00E-07	0.35	9.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	181	248	1.60E-05	-0.17	0.4		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	ж	0261	2062	1.80E-16	80.0	0.23		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2116	2163	0.0013	-0.25	0.05		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	æ	2178	2264	1.30E-11	-0.11	0.1		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	æ	2307	2371	5.40E-07	0.17	0.75		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2414	2458	0.0009	-0.14	0.58		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2620	2683	1.60E-05	0	90.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2692	2789	3.60E-19	0.56			N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	B	2942	3003	0.0036	0.42	0.92		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	æ	3044	3105	5.40E-06	0.56	0.42		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	æ	3146	3191	0.00036	-0.2	0.17		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
808	Inci	æ	3225	3313	1.80E-10	19.0	0.49		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	3354	3418	1.80E-08	0.87	66'0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	715	812	1.80E-19	0.15	96'0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	862	917	5.40E-05	-0.4	0.71		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	lnci	В	932	1022	1.80E-05	0.36	0.63		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Incj	٧	1039	1234	3.60E-36	0.45			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
808	lncj	∢	1147	1338	5.40E-33	0.34	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
808	lncj	٧	1270	1440	1.80E-22	0.3	0.54		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL

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PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN																			
Coumpound		N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN: CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN, CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN: CHAIN: A;											
SeqFold score																	120.25				
PMF score		0.99	-	0.21	_	0.74	96.0	0.76	0.33	98.0	-	- ·	0.75	0.83	0.3	-		-	0.99	_	0.58
Verify Score		0.29	0.03	0.05	0.45	-0.05	0.31	0.42	0.11	0.48	0.17	0.27	-0.1	0.14	0.25	0.29		0.28	0.31	0.62	-0.03
PSI- BLAST		3.60E-53	1.80E-34	5.40E-24	3.60E-33	3.60E-32	7.20E-27	3.60E-28	7.20E-55	3.60E-30	5.40E-52	5.40E-36	7.20E-32	1.60E-41	3.60E-27	1.30E-32	5.40E-63	5.40E-63	3.60E-38	1.80E-29	1.40E-31
End		1547	1652	354	1750	1861	0961	2062	2163	2264	2371	2473	2577	2683	450	2789	2897	2898	3003	3105	3196
Start AA		1351	1458	155	1562	1991	1782	1898	0261	2079	2178	2300	2407	2488	256	2593	1692	2693	2825	2913	3039
Chain TO		A	¥	V V	¥	A	V	٧	A	٧	¥	V	¥	A	Y	Ą	¥	٧	¥	V	A
PDB TD		Incj	Incj	lncj	Incj	Incj	Incj	lncj	lncj	lncj	1ncj	Incj	lncj	l ncj	1ncj	Incj	1 n cj	Incj	Incj	Incj	Incj
SEQ ID		808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	COMPLEX (BLOOD COAGULATION/INHBITOR) CHRISTMAS FACTOR; COMPLEX, INHIBITOR, HEMOPHILIA/EGF, BLOOD COAGULI ATION 2 PLASMA	SERINE PROTEASE, CALCIUM- BINDING, HYDROLASE, 3 GLYCOPROTEIN	SERINE PROTEASE FVIIA, FVIIA, BLOOD COAGULATION, SERINE PROTEASE	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE										
Coumpound	N-CADHERIN; CHAIN: A;	FACTOR IXA, CHAIN: C, L., D-PHE-PRO-ARG; CHAIN: I,		COAGULATION FACTOR VIIA (LIGHT CHAIN), CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL										
SeqFold score															
PMF score	0.63	0.99	_	0.39	0.21	0.35	66.0	0.18	_	0.88	_	-0.19		-0.19	0.57
Verify score	0.36	0.48	0.63	0.52	-0.28	0.3	0.37	0.02	0.23	0.34	0.39	0.01		0.2	89.0
PSI- BLAST	1.80E-34	7.20E-51	3.60E-32	3.60E-13	1.80E-57	1.80E-34	9.00E-30	1.80E-25	7.20E-62	1.80E-34	1.10E-34	1.40E-10		1.30E-11	1.20E-21
End	3313	3418	3523	3621	248	556	662	812	917	1022	1129	4312	-	4001	4033
Start AA	3120	3225	3346	3433	39	390	467	571	717	827	932	4224		3943	3951
Chain ID	V	A	A	A	<	<	¥.	4	<	¥	∢	1			ب
PDB ID	lncj	1ncj	Incj	lncj	lncj	Ipfx		1qfk	lqfk						
SEQ NO.	808	808	808	808	808	808	808	808	808	808	808	808		808	808

PDB annotation		SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE	COAGULATION FACTOR SERINE PROTEINASE, BLOOD COAGULATION, COAGULATION FACTOR	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
Coumpound	INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR: CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR IX; CHAIN: A; COAGULATION FACTOR IX; CHAIN: B;	EPITHELIAL CADHERIN; CHAIN: NULL;							
SeqFold score												,
PMF score		-0.17	-0.17	-0.19	0.27	0.93	0.45	0.62	0.13	0.04	0.63	0.07
Verify		0	0.37	0.03	0.24	-0.06	0.04	0.54	0.28	0.37	0.45	0.35
PSI- BLAST		5.40E-14	7.20E-13	7.80E-14	7.80E-20	1.60E-07	1.30E-17	1.30E-10	0.0013	1.80E-19	1.30E-08	2.60E-07
End		4202	4344	4001	1133	1133	1238	1334	1342	1444	1549	226
Start AA		4121	4267	3951	1041	9901	1145	1249	1279	1350	1455	155
Chain 10		٦	-1	а .								
PDB TD		1qfk	lqfk	if:	lsuh	Isuh	Isuh	Isuh	1suh	lsuh	1suh	lsuh
SEQ B NO:		808	808	808	808	808	808	808	808	808	808	808

	ELL	ELL	ELL	ELL	ELL	CELL	CELL							
	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL. ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN. CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL			
PDB annotation	CELL ADHESION UVOMORULN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UYOMORULM CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN CADHERIN, CALCIUM BINDING, ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING,
РДВ яп	SION U	SION U	SION U	SION U	SION U	SION U CALCI	SION U	SION U	SION U	SION L	SION L	SION L	SION L	CALCI
	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	CELL ADHE CADHERIN, ADHESION	L ADHE HERIN,
	CELL CADI ADHI	CELL	CELL	CELL CADI ADH	CELI CAD ADH	CELI CAD ADH	CELI CAD ADH	CELI	CELL	CELL	CELL ADH	AD SEL	CEL	CEL
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punod	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN: CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;							
Coumpound	EPITHELIAL C. CHAIN: NULL;	EPITHELIAL C. CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;	EPITHELIAL C CHAIN: NULL;
	EPITHE CHAIN:	EPITHE CHAIN:	CHAIN	EPITHE	CHAIN	EPITHE CHAIN	EPITHE CHAIN	CHAIN	CHAIN	CHAIN	CHAIN	EPITHI CHAIN	EPITHI	CHAIN
SeqFold score														
								_						,,
PMF	0.31	0.13	0.68	0.75	0.48	0.45	0.13	0.09	0.4	0.77	0.94	0.58	0.25	0.15
Verify score	0.41	-0.2	0.07	0.17	-0.06	-0.18	-0.41	0.42	-0.37	0.44	0.35	0.03	-0.04	-0.37
PSI- BLAST	1.00E-12	9.00E-09	1.30E-12	3.60E-06	1.30E-14	3.60E-05	1.80E-06	2.60E-15	1.30E-15	3.90E-20	3.60E-09	3.90E-05	3.60E-05	1.80E-12
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End	1650	1656	1754	1754	1867	1868	252	2163	2268	2375	2375	2475	2477	2581
Start AA	1570	1589	1991	1690	7771	1800	182	2084	2178	2280	2306	2395	2414	2488
Chain ID														
PDB ID	1suh	lsuh	lsuh	Isuh	lsuh	Isuh	lsuh	1suh	Isuh	lsuh	1suh	1suh	lsuh	1suh
SEQ ID	808	808	808	808	808	808	808	808	808	808	808	808	808	808

SEQ ID NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
										ADHESION
808	Isuh		2489	2581	1.30E-13	0.16	86.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2591	2681	2.60E-10	0.02	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2692	2793	3.60E-23	0.42	0.99		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2806	2902	2.60E-21	0.43	0.87		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	l suh		2831	2902	3.60E-09	0.01	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2914	3007	1.20E-14	0.37	0.93		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN. CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Suh		2941	3007	1.80E-06	-0.07	0.86		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		3026	3109	2.60E-21	0.36	0.89		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh	,	3046	3109	3.60E-06	0.4	0.72		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		3120	3185	0.0013	0.1	0.28		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		3120	3213	3.90E-12	0.43	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELI, ADHESION
808	Isuh		3225	3317	1.10E-14	9.0	0.82		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		3330	3422	3.90E-21	0.58	0.98		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		3355	3422	1.30E-09	99.0	0.93	х	EPITHELIAL CADHERIN,	CELL ADHESION UVOMORULIN;

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	2	<u> </u>	V V	¥ ¥	BLAST	score	score	score		
†									CHAIN: NULL;	CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		3435	3524	6.50E-11	0.56	0.65		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		363	454	0.0001	80.0	0.24		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		406	454	1.60E-05	-0.6	0.19		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		467	260	1.30E-15	-0.03	0.19		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERM, CALCIUM BINDING, CELL ADHESION
808	Isuh		590	658	3.90E-05	0.39	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN. CALCIUM BINDING, CELL ADHESION
808	Isuh		716	816	1.40E-23	-0.21	0.95		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		826	921	1.00E-17	0.4	0.72		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		854	921	5.40E-07	0.17	0.45		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		930	1026	5.20E-13	0.37	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	ltpg		3949	4026	1.30E-18	0.21	0.59		T-PLASMINOGEN ACTIVATOR FI-G; ITPG 7 CHAIN: NULL; ITPG 8	PLASMINOGEN ACTIVATION
808	lxka		4267	4348	5.40E-12	0.15	-0.19		BLOOD COAGULATION FACTOR XA; CHAIN: L, C;	BLOOD COAGULATION FACTOR STUART FACTOR, BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR LIKE DOMAIN
808	9wga	¥.	4166	4337	3.60E-10	0.05	-0.2		LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	
808	laut	1	3857	3947	6.50E-19	0.55	-0.12		ACTIVATED PROTEIN C,	COMPLEX (BLOOD

PDB annotation	COAGULATION/INHIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/INHIBITOR)	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX
Соитроипа	CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG- CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C.	BLOOD COAGULATION FACTOR VIIA; CHAIN: I, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG- CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I: DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D: PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y:
SeqFold score						
PMF score		0.07	0.07	0.15	0.05	-0.07
Verify score		0.09	0.1	-0.08	0.48	0.12
PSI- BLAST		1.40E-12	3.60E-14	1.40E-12	3.90E-21	3.60E-14
End		3897	3976	3897	3947	3976
Start AA		3831	3900	3831	3858	3900
Chain ID		-	ı	₽	<u> </u>	
PDB ID		ldan	ldan	ldva	ldva	ldva
SEQ ID NO:		808	808	808	808	808

PDB annotation	A L,	HYDROLASE/HYDROLASE INHIBITOR N: H, PROTEIN-PEPTIDE COMPLEX IA I. L, AIN: AIN:		4, B; CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	A, B; CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	A, B, CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	A, B, CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	t
Coumpound	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;	
SeqFold								
PMF	-0.15	-0.19	_	96:0	0.93	0.89	0.65	
Verify score	0.05	0.25	0.35	0.16	0.23	0.3	0.03	
PSI- BLAST	3.60E-12	1.30E-13	1.10E-33	9.00E-30	1.60E-20	3.60E-54	7.20E-32	2 (07)
End	4168	4259	1234	1338	1440	1547	1652	1760
Start AA	4078	4170	1026	1171	1279	1352	1455	1 6 00
Chain ID	٦	٦	V	∢	V	∢	⋖	
PDB ID	ldva	Idva	1edh	ledh	ledh	ledh	ledh	4
SEQ TO	808	808	808	808	808	808	808	808

									
PDB annotation	AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADI2; CADHERIN, CELL
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHÁIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;				
SeqFold score									
PMF score		0.25	66.0	0.07	0.94	0.34	0.92	_	-1.41
Verify score		-0.18	0.21	0.17	0.3	0.13	0.19	8 0.0	0.37
PSI- BLAST		3.60E-28	1.80E-27	5.40E-20	1.80E-30	1.80E-50	9.00E-29	1.80E-38	1.80E-32
End		0981	1960	354	2062	2163	2264	2371	2473
Start AA		1690	1780	182	1898	1975	2104	2178	2306
Chain ID		∢	V	¥	¥	4	<	<	¥
PDB ID		ledh	ledh	ledh	ledh	1edh	ledh	ledh	ledh
SEQ D		808	808	808	808	808	808	808	808

PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD 12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECADI2; CADHERIN, CELL	AUHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	EDITITE 141 CAPHERIN DOMAINS 1	AND TOADIS CADIEBRI CELL	AND 2, ECADIZ; CADRERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12: CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	NIGHOUS CAICUIS
Coumpound		E-CADHERIN; CHAIN: A, B;		E-CADHERIN, CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B:				a ratio inaditato a	E-CADRIERIN; CHAIN: A, B;				E-CADHERIN: CHAIN: A. B.					E-CADHERIN; CHAIN: A, B;					E-CADHERIN; CHAIN: A, B;				
SeqFold score																																			
PMF score		9.0	ļ	0.59				86.0				_				,	 				0.89					0.77					_				
Verify score		0.28		0.15				0.53	_			0.47				2	71.0				0.31					0.5					0.33				_
PSI- BLAST		1.80E-27		1.80E-28				1.40E-35				1.80E-16				7 700 63	3.00E-33				5.40E-30					3.60E-27					1.10E-57				
End		3191		3313				3418				3523				97.0	740				556					799					917				
Start		3045		3147				3225				3355				0,0	٧٠_	-			401					465					718				
Chain ID		V		٧				4				٧				4	٤				V					٨					∀				
PDB ID		1edh		ledh				ledh				Icdh				14.5			_	-	ledh					ledh					ledh			-	
SEQ ID NO:		808		808				808				808				000	000	_			808					808					808				_

SEQ ID NO:	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
808	ledh	<	854	1022	1.80E-33	0.14			E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	ledh	4	959	1129	1.80E-30	0.24	0.52	·	E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
808	lem n		3860	3931	3.60E-16	90.0	0.48		FIBRILLIN, CHAIN: NULL;	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-I FRAGMENT, MATRIX PROTEIN
808	lem n		4126	4217	9.00E-14	0.02	-0.18		FIBRILLIN; CHAIN: NULL;	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN
808	lfak	J	3900	3976	3.60E-14	0.12	-0.01		BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; SLI5; CHAIN: I;	BLOOD CLOTTING COMPLEX(SERNE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING
808	Ifak	1	4170	4259	1.30E-13	0.2	-0.19		BLOOD COAGULATION FACTOR VIIA, CHAIN. L, BLOOD COAGULATION FACTOR VIIA, CHAIN. H, SOLUBLE TISSUE FACTOR, CHAIN: T, 5L15, CHAIN: I,	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), PRODEASE/COFACTOR/LIGAND),
808	1klo		4074	4228	1.80E-17	0.07	-0.2		LAMININ; CHAIN: NULL;	GLYCOPROTEIN GLYCOPROTEIN

PDB Chain Start End ID AA AA B	Start End AA AA	End AA			PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
\top	4262 1.80E-18 0.07	4262 1.80E-18 0.07	1.80E-18 0.07	0.07	T	9	2		LAMININ; CHAIN: NULL;	GLYCOPROTEIN GLYCOPROTEIN
Incg 1064 1127 1.60E-05 0.12 0.63	1127 1.60E-05 0.12	1127 1.60E-05 0.12	1.60E-05 0.12	0.12		9.0	_		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1232 5.40E-05 0.15 0.87	1232 5.40E-05 0.15	1232 5.40E-05 0.15	5.40E-05 0.15	0.15		0.8	7		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1350 1439 3.60E-17 0.34 0.3	1439 3.60E-17 0.34	1439 3.60E-17 0.34	3.60E-17 0.34	0.34		0.3			N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1455 1546 5.40E-05 0 0	1546 5.40E-05 0	1546 5.40E-05 0	5.40E-05 0	0			0.72		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 155 229 3.60E-05 0.15 0	229 3.60E-05 0.15	229 3.60E-05 0.15	3.60E-05 0.15	0.15	<u> </u>		60.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1599 1651 9.00E-06 -0.45	1651 9.00E-06	1651 9.00E-06	9.00E-06		-0.45		0.04		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1667 1748 3.60E-06 0.35	1748 3.60E-06	1748 3.60E-06	3.60E-06		0.35		0.31		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1975 2061 1.80E-15 -0.06	2061 1.80E-15	2061 1.80E-15	1.80E-15		90.0-		0.1		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 2079 2161 0.00036 0.31	2161 0.00036	2161 0.00036	0.00036		0.31		0.29		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 2180 2262 9.00E-07 0.11	2262 9.00E-07	2262 9.00E-07	9.00E-07		0.11		10.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 2304 2369 1.80E-06 0.43	2369 1.80E-06	2369 1.80E-06	1.80E-06		0.43		0.64		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 2411 2458 0.0045 -0.23	1 2458 0.0045	1 2458 0.0045	0.0045		-0.23		0.22		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 2592 2681 0.00014 0.41	2681 0.00014	2681 0.00014	0.00014		0.41		0.25		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 2692 2788 3.60E-20 0.44	2788 3.60E-20	2788 3.60E-20	3.60E-20		0.44		2.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
	3106 9.00E-07	3106 9.00E-07	9.00E-07		0.45		0.51		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 3120 3191 0.00018 0.06	3191 0.00018	3191 0.00018	0.00018		90.0		0.34		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 3225 3312 1.10E-12 0.28	3312 1.10E-12	3312 1.10E-12	1.10E-12		0.28		0.45		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
lncg 716 811 1.10E-21 0.16	811 1.10E-21	811 1.10E-21	1.10E-21		0.16		8.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 852 900 0.00036 -0.19	9000 0.00036	9000 0.00036	0.00036		-0.19		0.53		N-CADHERIN, INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 932 1021 1.60E-05 0.55	1021 1.60E-05	1021 1.60E-05	1.60E-05		0.55	·	0.46		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN

SEQ NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
808	lnci	В	1065	1129	3.60E-06	90.0	0.78		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1178	1234	1.80E-05	-0.16	6.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1350	1440	3.60E-16	0.53	90.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1481	1547	5.40E-05	0.21	0.16		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1991	1750	1.40E-06	0.26	0.64		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	181	248	5.40E-05	-0.17	0.4		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1912	1960	0.0045	-0.5	0.27		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	1975	2062	1.80E-14	0.24	0.07		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2180	2264	3.60E-06	-0.4	0.3		N-CADHERIN, INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2307	1782	5.40E-07	6E'0	9.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2412	2458	0.0079	-0.03	0.49		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2692	5789	1.80E-19	0.56	1		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2832	2898	1.80E-06	-0.01	0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
808	1nci	В	2951	3003	0.0045	-0.24	0.78		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	3041	3105	1.10E-06	0.52	99.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	3146	3191	0.0013	-0.2	0.17		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	æ	3225	3313	1.40E-10	0.31	0.17		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
808	Inci	В	716	812	5.40E-20	10.0	0.88		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
808	Inci	В	853	917	7.20E-05	0.26	0.84		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	932	1022	7.20E-06	0.4	0.59		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Incj	A	1039	1234	1.10E-35	0.39	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL

PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN		CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN						CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN		CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN			CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	
Coumpound		N-CADHERIN: CHAIN: A;	N-CADHERIN: CHAIN: A;	N-CADHERIN, CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN, CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN, CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;
SeqFold score																			120.25		
PMF			0.16	66.0	0.88	_	0.84	0.23	-1	6.0	0.45	68.0	-		8.0	9.0	0.51			_	0.99
Verify score		0.53	0.1	0.28	0.02	0.46	0.2	0.08	0.31	0.25	0.02	0.23	0.25	0.32	-0.03	90.0	0.28	0.23		0.28	0.28
PSI- BLAST		1.40E-31	1.80E-23	1.60E-58	3.60E-35	1.10E-33	7.20E-33	3.60E-22	9.00E-27	3.60E-31	1.60E-56	9.00E-31	3.60E-40	3.60E-35	9.00E-31	3.60E-40	1.30E-26	1.40E-31	3.60E-63	3.60E-63	9.00E-36
End		1338	1440	1547	1652	1750	1861	354	1960	2062	2163	2264	2371	2473	2577	2683	450	2789	2897	2898	3003
Start		1144	1251	1351	1455	1562	1991	173	1793	1898	1975	2079	2180	2300	2407	2488	256	2592	2691	2693	2825
Chain ID		V	V	ĕ.	<	¥	¥	¥	<	V	<	∢	<	<	⋖	∢	<	⋖	∢	V	V
PDB ID		Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	Incj	1ncj	Incj	Incj	Incj	lncj	Incj
SEQ ID		808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808	808

	Chain Start End	End		PSI-		Verify	PMF	SeqFold	Coumpound	PDB annotation
ID AA AA BLAST score	ID AA AA BLAST score	AA BLAST score	BLAST score	score		-1	score	score		I lao idamoda wowe.
Incj A 2915 3105 9.00E-30 0.35 1	A 2915 3105 9.00E-30	3105 9.00E-30	105 9.00E-30		0.35 1	-	}		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3039 3196 1.30E-30 0.16 0.8	A 3039 3196 1.30E-30 0.16	3196 1.30E-30 0.16	196 1.30E-30 0.16	0.16		0.8			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3141 3313 1.30E-30 0.35 0.82	A 3141 3313 1.30E-30 0.35	3313 1.30E-30 0.35	313 1.30E-30 0.35	0.35		0.8	2)		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3225 3418 1.80E-39 0.66 1	A 3225 3418 1.80E-39	3418 1.80E-39	1.80E-39		0.66	-			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3349 3523 1.80E-17 0.6 1	A 3349 3523 1.80E-17	3523 1.80E-17	1.80E-17	<u> </u>	0.6	-			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3473 3621 5.40E-10 0.35 -0.	A 3473 3621 5.40E-10 0.35	3621 5.40E-10 0.35	5.40E-10 0.35	0.35		Ó.	-0.03		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 39 248 1.10E-59 -0.28 0.21	A 39 248 1.10E-59 -0.28	248 1.10E-59 -0.28	1.10E-59 -0.28	-0.28		0.2	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 390 556 1.80E-35 0.08 0.36	A 390 556 1.80E-35 0.08	556 1.80E-35 0.08	1.80E-35 0.08	0.08	 	0.3	9		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 467 662 7.20E-28 0.36 0.96	A 467 662 7.20E-28 0.36	662 7.20E-28 0.36	7.20E-28 0.36	0.36	<u> </u>	0.9	\ 		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 573 812 1.80E-25 -0.13 0.01	A 573 812 1.80E-25 -0.13	812 1.80E-25 -0.13	1.80E-25 -0.13	-0.13	1	0.0	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 717 917 1.10E-63 0.39 1	A 717 917 1.10E-63	917 1.10E-63	1.10E-63	ļ -	0.39 1	-			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 827 1022 1.80E-34 0.35 0.88	A 827 1022 1.80E-34 0.35	1022 1.80E-34 0.35	1.80E-34 0.35	0.35	<u> </u>	0.88			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 932 1129 3.60E-34 0.48 1	A 932 1129 3.60E-34	1129 3.60E-34	3.60E-34		0.48	_			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
1qfk L 3865 3947 1.20E-21 0.68 0.57	L 3865 3947 1.20E-21 0.68	3947 1.20E-21 0.68	1.20E-21 0.68	89.0		0.5	_		COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE
									VIIA (HEAVI CHAIN), CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	
1qfk L 3904 3976 3.60E-13 0.01 -0.08	L 3904 3976 3.60E-13 0.01	3976 3.60E-13 0.01	3.60E-13 0.01	3.60E-13 0.01		-0.0			COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN:	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE
									VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	
1qfk L 4082 4168 7.20E-11 0.08 -0.15	L 4082 4168 7.20E-11 0.08	4168 7.20E-11 0.08	7.20E-11 0.08	7.20E-11 0.08		o O	100		COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR	SERINE PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE

SEQ ID NO:	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
									VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	
808	1suh		1026	1133	7.20E-08	0.33	0.13		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		1041	1133	7.80E-20	0.24	0.27		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1145	1238	1.30E-17	0.04	0.45		EPITHELIAL CADHERIN; CHAIN: NULL:	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1171	1238	7.20E-08	-0.03	0.4		EPITHELIAL CADHERIN: CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1 suh		1249	1334	1.30E-10	0.54	0.62		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1350	1444	1.80E-21	0.3	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1455	1549	1.30E-08	0.45	0.63		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1455	1551	5.40E-07	0.14	0.11		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		155	226	2.60E-07	0.35	0.07		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1570	0591	1.00E-12	0.41	0.31		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1589	9591	1.40E-08	0.2	60.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1 suh		1667	1754	1.30E-12	0.07	89.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		1690	1754	3.60E-06	90.0	99.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL

SEQ	PDB ID	Chain	Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
NO:										
										ADHESION
808	1suh		7771	1867	1.30E-14	90:0-	0.48		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1780	1868	3.60E-05	-0.24	0.39		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		182	252	5.40E-07	-0.41	0.13		EPITHELIAL CADHERIN; CHAIN: NULL:	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		8681	1964	1.60E-05	-0.1	0.18		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1975	2066	7.20E-18	0.17	1202.08		EPITHELIAL CADHERÎN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2084	2163	2.60E-15	0.42	0.09		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2104	2167	3.60E-07	0.05	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2178	2268	1.30E-14	-0.23	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2178	2268	3.60E-10	-0.36	0.46		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2280	2375	3.90E-20	0.44	0.77		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2306	2375	1.10E-08	0.4	0.94		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2395	2475	3.90E-05	0.03	0.58		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION
808	l suh		2414	2477	0.00011	-0.37	90.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2488	2581	3.60E-12	-0.2	0.03		EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

1	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	:	1		5		31000	31036	3000		
									CHAIN: NULL;	CADHERIN, CALCIUM BINDING, CELL ADHESION
	1suh		2489	2581	1.30E-13	0.16	86.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	Isuh		2591	2681	2.60E-10	0.02	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	Isuh		2692	2793	1.40E-23	0.51	0.99		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	lsuh		2806	2902	2.60E-21	0.43	0.87		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	lsuh		2831	2902	1.40E-08	-0.01	0.53		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	lsuh		2914	3007	1.20E-14	0.37	0.93		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	l suh		2941	3007	1.80E-05	-0.07	0.86		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL, ADHESION
	1suh		3026	3109	2.60E-21	0.36	0.89		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERN, CALCIUM BINDING, CELL ADHESION
	1suh		3045	3109	3.60E-06	0.33	0.55		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	1 suh		3120	3213	3.90E-12	0.43	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	1 suh		3225	3317	1.30E-13	9.0	0.82		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERN, CALCIUM BINDING, CELL ADHESION
	l suh		3330	3422	3.90E-21	0.58	86.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
	Isuh		3355	3422	5.40E-05	9.0	0.94		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION

	CELL		 ЭМТН									
PDB annotation	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESTON UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESTON	CELL ADHESION UVOMORULIN; CADHERIN. CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	PLASMINOGEN ACTIVATION	BLOOD COAGULATION FACTOR STUART FACTOR, BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH
	CEL	PLA	STU, COA PRO									
Coumpound	EPITHELIAL CADHERIN; CHAIN: NULL;	T-PLASMINOGEN ACTIVATOR F1-G; 17PG 7 CHAIN: NULL: 17PG 8	BLOOD COAGULATION FACTOR XA; CHAIN: L, C,									
SeqFold score									:			
PMF score	0.65	0.24	0.3	0.19	0.57	0.95	0.72	0.47	0.57	0.18	0.59	0.1
Verify score	0.56	80.0	-0.21	-0.03	0.39	-0.21	0.4	0.27	0.37	0.2	0.21	-0.26
PSI- BLAST	6.50E-11	0.0001	9.00E-06	1.30E-15	3.90E-05	7.20E-25	1.00E-17	1.30E-07	5.20E-13	9.00E-07	1.30E-18	7.20E-11
End	3524	454	454	260	859	816	921	921	1026	1026	3940	3994
Start	3435	363	401	467	290	716	826	854	930	656	3863	3904
Chain ID												٦
PDB ID	lsuh	Isuh	lsuh	1suh	1suh	l suh	Isuh	1suh	1suh	1 suh	ltpg	lxka
SEQ ID	.08 808	808	808	808	808	808	808	808	808	808	808	808

PDB annotation	SERINE PROTEINASE COAGULATION FACTOR II; FETOMODULIN, TM, CD141 ANTIGEN; EGR-CMK SERINE PROTEINASE, EGF-LIKE DOMAINS, ANTICOAGULANT COMPLEX, 2 ANTIFIBRINOLYTIC COMPLEX	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL
Coumpound	THROMBIN LIGHT CHAIN; CHAIN: A, B, C, D; THROMBIN HEAVY CHAIN; CHAIN: M, N, O, P; THROMBOMODULIN; CHAIN: I, K, L, THROMBIN INHIBITOR L-GLU-L-GLY-L- ARM; CHAIN: E, F, G, H;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A. B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold score								
PMF	-0.19	-	1202.08	6.0	0.82	-1.41	96.0	66.0
Verify score	0.06	0.26	0.22	0.45	0.27	0.1	0.21	0.19
PSI- BLAST	1.30E-11	3.60E-33	1.30E-32	3.60E-21	1.60E-49	1.30E-32	3.60E-33	1.60E-28
End	4336	1234	1338	1440	1547	1652	1750	1860
Start	4232	1066	1171	1279	1352	1460	1589	0691
Chain ID	-	<	V	<	<	⋖	<	¥
PDB ro	Idx5	ledh	1edh	ledh	ledh	1edh	1edh	ledh
SEQ ID	608	808	808	808	808	808	808	808

т		· · · · · · · · · · · · · · · · · · ·			- 1	γ			
PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound		E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;						
SeqFold score									
PMF		_	0	96.0	0.51	0.78	_		0.86
Verify score		0.32	0	0.26	0.3	0.24	0.18	0.32	0.07
PSI- BLAST		3.60E-26	1.60E-20	3.60E-29	1.10E-50	1.30E-28	3.60E-48	1.60E-35	1.80E-29
End		0961	354	2062	2163	2264	2371	2473	2577
Start		1800	182	1898	1975	2104	2178	2306	2414
Chain ID		<	<	<	K	4	<	₹	⋖
PDB		1edh	ledh	ledh	ledh	ledh	ledh	ledh	ledh
SEQ	Ž.	808	808	808	809	608	808	808	608

PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I	AND 2, ECADI2; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I AND 2. ECADI2; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN FPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12, CADHERIN, CELL	ADHESION PROTEIN, CALCIUM RINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELLA CADHERIN DOMAINS 1	AND 2 ECADIS: CADHERIN. CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHEI 141 CADHERIN DOMAINS I	AND 3 ECADIS CADHERIN CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 4, ECADI2, CADRENIN, CELE ADHESION PROTEIN, CALCIUM	BINDING PROTEIN
Coumpound		E-CADHERIN; CHAIN: A, B;		E-CADHERIN; CHAIN: A, B;			E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;			F-CADHERIN CHAIN: A. B.					E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;			
SeqFold score							120.62	_																						
PMF		0.93		96.0						***	I			-	•				0.94				0.53				0.64			
Verify score		0		0.1							0.47			0.27	3				0.31				0.38				91.0			
PSI- BLAST		1.80E-38		5.40E-32			1.80E-57				1.80E-57			7 20E-35	CC-707.1				5.40E-29				7.20E-25				5.40E-28			
End		2683		2789			2895				2898			3003	COAC				3105				450				3197			
Start		2488		2619			2692				2693			2021	1607				2941				596				3046			
Chain ID		<		<			V				4				ζ				V				4				<			_
PDB ID		ledh		ledh			ledh				ledh			460	1031				ledh				ledh				ledh			
SEQ ID		808		608			808				608			000	600				808				809				809			

PDB Chain Start End ID AA AA	Start			PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
ledh A 3120 3313 I.80E-32 0.57	3313 1.80E-32	1.80E-32		0.57		-1.41		E-CADHERIN, CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
ledh A 3225 3418 1.80E-48 0.45	3418 1.80E.48	1.80E-48		0.45		0.98		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12: CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
1edh A 3355 3523 3.60E-30 0.66	3523 3.60E-30	3523 3.60E-30		99.0		-		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
1edh A 39 248 1.80E-51 0.12	248 1.80E-51	1.80E-51		0.12		0.3		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
556	556 1.10E-29	1.10E-29		0.29		0.83		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
ledh A 464 662 3.60E-29 0.17	662 3.60E-29	3.60E-29		0.17		0.82		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
ledh A 591 812 5.40E-22 -0.05	812 5.40E-22	5.40E-22	<u> </u>	-0.05		0.27		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
1edh A 718 917 1.10E-55 0.33	917 1.10E-55	1.10E-55		0.33		_		E-CADHERIN, CHAIN: A, B,	CELL ADHESION PROTEIN EPITHELIAL CADHERN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Tedh A 854 1022 1.10E-32 0.31	1022 1.10E-32	1.10E-32	1.10E-32	0.3				E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN

1 1 1 1 1 1 1 1 1 1	SEQ U	PDB	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
1edh A 959 1129 9,00E-32 0,11 0,69 E-CADHERIN, CHAIN; A, B, B, C 1,80E-16 0,71 0,99 FIBRILLIN; CHAIN; NULL; A 1,10E 1,30E-11 1,03 0,88 P-SELECTIN; CHAIN; NULL; A 1,10E 1,3954 4056 1,10E-13 -0,1 0,27 LAMININ; CHAIN; NULL; A 1,10E 1,3954 4056 1,10E-13 -0,1 0,27 LAMININ; CHAIN; NULL; A 1,10E 1,	sl s										EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
16m 3946 4022 1.80E-16 0.71 0.99 FIBRILLIN, CHAIN: NULL; N N N N N N N N N	808	1edh	<	959	1129	9.00E-32	0.11	0.69		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
145b 3949 3988 1.30E-11 1.03 0.88 P-SELECTIN, CHAIN: NULL; Colored 1450 1.10E-13 -0.1 0.27 LAMININ; CHAIN: NULL; Colored 1420 4342 3.60E-17 0.23 LAMININ; CHAIN: NULL; Colored 1.27 9.00E-06 0.12 0.29 N-CADHERIN: INCG 3 Colored 1.167 1.232 0.00014 0.13 0.69 N-CADHERIN: INCG 3 Colored 1.167 1.232 0.00014 0.13 0.69 N-CADHERIN: INCG 3 Colored 1.168 1.40E-14 0.01 0.29 N-CADHERIN: INCG 3 Colored 1.168 1.169	6	n n		3946	4022	1.80E-16	0.71	0.99		FIBRILLIN; CHAIN: NULL;	MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN
Iklo 3924 4050 1.10E-13 -0.1 0.27 LAMININ, CHAIN: NULL; O Iklo 3954 4075 5.40E-21 0.07 -0.03 LAMININ, CHAIN: NULL; O Iklo 4201 4342 3.60E-17 0.23 -0.2 LAMININ; CHAIN: NULL; O Iklo 1062 1127 9.00E-06 0.12 0.29 N-CADHERIN; INCG 3 Incg 1167 1232 0.00014 0.13 0.69 N-CADHERIN; INCG 3 Incg 1350 1439 1.40E-14 0.01 0.29 N-CADHERIN; INCG 3 Incg 155 229 3.60E-05 0.15 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1970 2061 0.22 0.11 N-CADHERIN; INCG 3 Incg 1970 2061 0.22 0.1 N-CADHERIN; INCG 3 Incg 1079 2161 0.00036 0.28	6	1fsb		3949	3988	1.30E-11	1.03	0.88		P-SELECTIN; CHAIN: NULL;	CELL ADHESION PROTEIN EGF-LIKE DOMAIN, CELL ADHESION PROTEIN, TRANSMEMBRANE, 2 GLYCOPROTEIN
Ikio 3954 4075 5,40E-21 0.07 -0.03 LAMININ; CHAIN: NULL; Ikio 4201 4342 3.60E-17 0.23 -0.2 LAMININ; CHAIN: NULL; Incg 1062 1127 9.00E-06 0.12 0.29 N-CADHERIN; INCG 3 Incg 1167 1232 0.00014 0.13 0.69 N-CADHERIN; INCG 3 Incg 1350 1439 1.40E-14 0.01 0.29 N-CADHERIN; INCG 3 Incg 155 229 3.60E-05 0.15 0.09 N-CADHERIN; INCG 3 Incg 1570 2061 3.60E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 1070 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3	6	1klo		3924	4050	1.10E-13	1.0-	0.27		LAMININ; CHAIN: NULL;	GLYCOPROTEIN GLYCOPROTEIN
Incg 1167 1232 0.006-06 0.12 0.29 N-CADHERIN; INCG 3 Incg 1167 1232 0.00014 0.13 0.69 N-CADHERIN; INCG 3 Incg 1350 1439 1.40E-14 0.01 0.29 N-CADHERIN; INCG 3 Incg 1480 1546 0.00018 0.46 0.1 N-CADHERIN; INCG 3 Incg 1559 1650 9.00E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	2 8	1klo		3954 4201	4342	3.60E-17	0.23	-0.03		LAMININ, CHAIN: NULL,	GLYCOPROTEIN GLYCOPROTEIN
Incg 1167 1232 0.00014 0.13 0.69 N-CADHERIN; INCG 3 Incg 1350 1439 1.40E-14 0.01 0.29 N-CADHERIN; INCG 3 Incg 155 229 3.60E-05 0.15 0.09 N-CADHERIN; INCG 3 Incg 1599 1650 9.00E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	60	Incg		1062	1127	9.00E-06	0.12	0.29		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1350 1439 1,40E-14 0.01 0.29 N-CADHERIN: INCG 3 Incg 1546 0.00018 0.46 0.1 N-CADHERIN: INCG 3 Incg 155 229 3.60E-05 0.15 0.09 N-CADHERIN; INCG 3 Incg 1599 1650 9.00E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 0.35 0.31 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	6	Incg		1167	1232	0.00014	0.13	69.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1480 1546 0.00018 0.46 0.1 N-CADHERIN; INCG 3 Incg 155 229 3.60E-05 0.15 0.09 N-CADHERIN; INCG 3 Incg 1599 1650 9.00E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 0.35 0.31 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	66	Incg		1350	1439	1.40E-14	0.01	0.29		N-CADHERIN: INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 155 229 3.60E-05 0.15 0.09 N-CADHERIN; INCG 3 Incg 1599 1650 9.00E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 0.35 0.31 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	66	Incg		1480	1546	0.00018	0.46	0.1		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1599 1650 9.00E-06 -0.13 0.09 N-CADHERIN; INCG 3 Incg 1667 1748 3.60E-06 0.35 0.31 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	66	Incg		155	229	3.60E-05	0.15	60.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1667 1748 3.60E-06 0.35 0.31 N-CADHERIN; INCG 3 Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	60	lncg		1599	1650	9.00E-06	-0.13	0.09		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 1970 2061 3.60E-17 0.22 0.1 N-CADHERIN; INCG 3 Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	60	Incg		1991	1748	3.60E-06	0.35	0.31		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
Incg 2079 2161 0.00036 0.28 0.11 N-CADHERIN; INCG 3	66	lncg		1970	2061	3.60E-17	0.22	0.1		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
	8	Incg		2079	2161	0.00036	0.28	0.11		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13

PDB annotation	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN 1NCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INC! 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN INCI 13	CELL ADHESION PROTEIN CADHERIN										
Coumpound	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3	N-CADHERIN; INCI 3	N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3	N-CADHERIN; INCI 3	N-CADHERIN; 1NCI 3
SeqFold score																					
PMF	0.1	0.74	0.43	0.57	0.7	0.39	0.51	0.34	0.36	8.0	0.53	0.64	0.78	96.0	0.22	9.65	9.0	0.4	0.23	0.05	0.1
Verify score	-0.08	0.41	-0.27	0.34	0.44	0.35	0.45	90.0	0.33	91.0	-0.19	0.38	90.0	90.0	0.43	-0.08	0.35	-0.17	0.08	-0.25	-0.11
PSI- BLAST	3.60E-12	1.80E-06	0.00036	1.60E-05	1.80E-19	0.00054	3.60E-06	1.80E-05	1.60E-11	7.20E-20	0.00018	5.40E-05	1.80E-06	5.40E-05	1.30E-13	0.00018	9.00E-07	1.60E-05	1.80E-16	0.0013	1.30E-11
End	2263	2370	2458	2681	2788	2988	3106	3191	3311	811	006	1003	1129	1234	1440	1547	1750	248	2062	2163	2264
Start	2178	2304	2411	2593	2692	2913	3039	3120	3225	716	852	932	1065	1172	1350	1491	1667	181	1970	2116	2178
Chain ID													В	В	æ	В	æ	æ	e B	B	В
PDB ID	Incg	Incg	lncg	lncg	lncg	1ncg	lncg	lncg	lncg	lncg	Incg	Incg	Inci	l nci	1nci	1nci	Inci	Inci	Inci	Inci	Inci
SEQ ID NO:	808	809	608	608	808	809	808	808	608	608	608	809	808	608	809	808	809	808	809	809	808

SEQ	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
ÿ.										1NCI 13
608	Inci	В	2307	2371	5.40E-07	0.17	0.75		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2414	2458	0.0000	-0.14	0.58		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
809	1nci	В	2620	2683	1.60E-05	0	90:0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	2692	2789	3.60E-19	0.56	_		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2942	3003	0.0036	0.42	0.92		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	3044	3105	5.40E-06	0.56	0.42		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
809	Inci	В	3146	3191	0.00036	-0.2	0.17		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	1nci	В	3225	3313	1.80E-10	19.0	0.49		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	3354	3418	1.80E-08	0.87	0.99		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
608	Inci	В	715	812	1.80E-19	0.15	96.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN
809	Inci	8	862	917	5.40E-05	-0.4	0.71		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
809	Inci	В	932	1022	1.80E-05	0.36	0.63		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
809	lncj	<	1039	1234	3.60E-36	0.45			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
809	Incj	4	1147	1338	5.40E-33	0.34	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
808	Incj	<	1270	1440	1.80E-22	0.3	0.54		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
809	1ncj	4	1351	1547	3.60E-53	0.29	0.99		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
809	Incj	4	1458	1652	1.80E-34	0.03			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
809	Incj	K	155	354	5.40E-24	0.05	0.21		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
808	Incj	4	1562	1750	3.60E-33	0.45	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
808	Incj	<	1991	1861	3.60E-32	-0.05	0.74		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN

1	1	F	DOT	1/0-:6.	DME	SearFold	Commound	PDB annotation
Chain Start ID AA		AA	BLAST	score	score	score		
1782		1960	7.20E-27	0.31	96.0		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
1898		2062	3.60E-28	0.42	0.76		N-CADHERIN: CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
1970		2163	7.20E-55	0.11	0.33		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2079		2264	3.60E-30	0.48	98.0		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2178		2371	5.40E-52	0.17	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2300		2473	5.40E-36	0.27			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2407		2577	7.20E-32	-0.1	0.75		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2488		2683	1.60E-41	0.14	0.83		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
256		450	3.60E-27	0.25	0.3		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2593	1	2789	1.30E-32	0.29	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2691	1	2897	5.40E-63			120.25	N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2693	1	2898	5.40E-63	0.28			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2825		3003	3.60E-38	0.31	66:0		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
2913	1	3105	1.80E-29	0.62	_		N-CADHERIN, CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
3039		3196	1.40E-31	-0.03	0.58		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
3120		3313	1.80E-34	0.36	0.63		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
3225	lio.	3418	7.20E-51	0.48	0.99		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
3346	1	3523	3.60E-32	0.63	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
3433	í	3621	3.60E-13	0.52	0.39		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
39	1	248	1.80E-57	-0.28	0.21		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
390		556	1.80E-34	0.3	0.35		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL

PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	COMPLEX (BLOOD COAGULATION/INHIBITOR) CHRISTMAS FACTOR; COMPLEX, NHIRITOR HEMOPHII IA/EGF	BLOOD COAGULATION, 2 PLASMA, SERINE PROTEASE, CALCIUM-	BINDING, HYDROLASE, 3 GLYCOPROTEIN	SERINE PROTEASE FVIIA;	BLOOD COAGOLATION, SERINE PROTEASE			SERINE PROTEASE FVIIA; FVIIA;	BLOUD COAGOLATION, SERINE PROTEASE			SERINE PROTEASE FVIIA; FVIIA;	BLOOD COAGOLATION, SENINE				SEKINE PROTEASE FVIIA; FVIIA;	PROTEASE					
Coumpound		N-CADHERIN; CHAIN: A;	FACTOR IXA; CHAIN: C, L,; D-PHE-PRO-ARG; CHAIN: I;			COAGULATION FACTOR	VIIA (LIGHT CHAIN); CHAIN:	VIIA (HEAVY CHAIN);	CHAIN: H; TRIPEPTIDYL NHIBITOR; CHAIN: C;	COAGULATION FACTOR	L; COAGULATION FACTOR	VIIA (HEAVY CHAIN);	CHAIN: H; I KIPEF I ID Y L INHIBITOR; CHAIN: C;	COAGULATION FACTOR	VIIA (LIGHI CHAIN); CHAIN:	VIIA (HEAVY CHAIN);	CHAIN: H; TRIPEPTIDYL	INHIBITOR; CHAIN: C;	COAGULATION FACTOR	L; COAGULATION FACTOR	VIIA (HEAVY CHAIN);				
SeqFold score																									
PMF		0.99	0.18	-	0.88	-	-0.19			-0.19		_		0.57				-0.17			_		-0.17		
Verify score		0.37	0.02	0.23	0.34	0.39	0.01			0.2				89.0		_		0					0.37		
PSI- BLAST		9.00E-30	1.80E-25	7.20E-62	1.80E-34	1.10E-34	1.40E-10			1.30E-11				1.20E-21				5.40E-14					7.20E-13		
End		662	812	917	1022	1129	4312			4001				4033				4202					4344		
Start		467	571	717	827	932	4224			3943				3951				4121					4267		
Chain ID		V	4	٧	Y	V	٦			L				L				1					-1		
PDB ID		lncj	Incj	Incj	Incj	Incj	1pfx			1qfk				1qfk	-			lqfk					198 		
SEQ ID		808	608	809	608	608	608			809				809				809					808		

								—-						
PDB annotation		COAGULATION FACTOR SERINE PROTEINASE, BLOOD COAGULATION, COAGULATION FACTOR	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING. CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN;
Coumpound	CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR IX; CHAIN: A; COAGULATION FACTOR IX; CHAIN: B;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL:	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN;								
SeqFold score														
PMF score		-0.19	0.27	0.93	0.45	0.62	0.13	0.04	0.63	0.07	0.31	0.13	0.68	0.75
Verify score		0.03	0.24	-0.06	0.04	0.54	0.28	0.37	0.45	0.35	0.41	-0.2	0.07	0.17
PSI- BLAST		7.80E-14	7.80E-20	1.60E-07	1.30E-17	1.30E-10	0.0013	1.80E-19	1.30E-08	2.60E-07	1.00E-12	9.00E-09	1.30E-12	3.60E-06
End		4001	1133	1133	1238	1334	1342	1444	1549	226	1650	1656	1754	1754
Start		3951	1041	1066	1145	1249	1279	1350	1455	155	1570	1589	1991	0691
Chain ID		В												
PDB ID		1rfn	Isuh	1suh	Isuh	lsuh	1suh	1suh	lsuh	1suh	1suh	Isuh	Isuh	Isuh
SEQ NO:		808	808	608	608	608	809	608	608	808	809	809	808	808

SEQ ID	PDB 10	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SegFold	Coumpound	PDB annotation
NO:										ADHESION
608	1suh		1777	1867	1.30E-14	-0.06	0.48		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		1800	1868	3.60E-05	81.0-	0.45		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		182	252	1.80E-06	-0.41	0.13		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2084	2163	2.60E-15	0.42	0.09		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2178	2268	1.30E-15	-0.37	0.4		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2280	2375	3.90E-20	0.44	0.77		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2306	2375	3.60E-09	0.35	0.94		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2395	2475	3.90E-05	0.03	0.58		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		2414	2477	3.60E-05	-0.04	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2488	2581	1.80E-12	-0.37	0.15		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2489	2581	1.30E-13	0.16	96.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		2591	2681	2.60E-10	0.02	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		2692	2793	3.60E-23	0.42	0.99		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2806	2902	2.60E-21	0.43	0.87		EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

SEQ NO:	PDB ID	Chain TO	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									CHAIN: NULL;	CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		2831	2902	3.60E-09	0.01	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2914	3007	1.20E-14	0.37	0.93		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2941	3007	1.80E-06	-0.07	0.86		EPITHELIAL CADHERIN; CHAIN: NULL:	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh .		3026	3109	2.60E-21	0.36	68.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		3046	3109	3.60E-06	0.4	0.72		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		3120	3185	0.0013	0.1	0.28		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		3120	3213	3.90E-12	0.43	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	lsuh		3225	3317	1.10E-14	9.0	0.82		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1 suh		3330	3422	3.90E-21	0.58	86.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		3355	3422	1.30E-09	0.66	0.93		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1 suh		3435	3524	6.50E-11	0.56	0.65		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		363	454	0.0001	0.08	0.24		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	l suh		406	454	1.60E-05	-0.6	0.19	·	EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION

SEQ EQ	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
808	1suh		467	999	1.30E-15	-0.03	61.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULM: CADHERIN. CALCIUM BINDING, CELL ADHESION
608	lsuh		290	658	3.90E-05	0.39	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		716	816	1.40E-23	-0.21	0.95		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULN; CADHERN, CALCIUM BINDING, CELL ADHESION
608	1suh		826	921	1.00E-17	0.4	0.72		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		854	921	5.40E-07	0.17	0.45		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	Isuh		930	1026	5.20E-13	0.37	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1tpg		3949	4026	1.30E-18	0.21	0.59		T-PLASMINOGEN ACTIVATOR F1-G; 1TPG 7 CHAIN: NULL; 1TPG 8	PLASMINOGEN ACTIVATION
808	Ixka	- -1	4267	4348	5.40E-12	0.15	-0.19		BLOOD COAGULATION FACTOR XA; CHAIN: L, C;	BLOOD COAGULATION FACTOR STUART FACTOR; BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR LIKE DOMAIN
608	9wga	٧	4166	4337	3.60E-10	0.05	-0.2		LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	
808	laut	1	3857	3947	6.50E-19	0.55	-0.12		ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	COMPLEX (BLOOD COAGULATION/INHIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD COAGULATION/INHIBITOR)
808	Idan	1	3831	3897	1.40E-12	0.09	0.07		BLOOD COAGULATION FACTOR VIIA; CHAIN: I, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE- ARG-	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COFACTOR/LIGAND)

PDB annotation			BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GIA FGF 3 COMPLEX (SERINE	PROTEASE/COFACTOR/LIGAND)		HYDROLASE/HYDROLASE INHIBITOR PROTEIN-PEPTIDE COMPLEX				HYDROL ASE/HYDROLASE INHIBITOR	PROTEIN-PEPTIDE COMPLEX					TIME OF ACTUADED A ACE INITION	PROTEIN-PEPTIDE COMPLEX						PROTEIN-PEPTIDE COMPLEX						PROTEIN-PEPTIDE COMPLEX	
Coumpound		CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T 11: D.PHF.PHF.	ARG-	(DFFRCMK) WITH CHAIN: C;	DES-GLA FACTOR VIIA	I; DES-GLA FACTOR VIIA	(LIGHT CHAIN); CHAIN: L,	M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN:	DES.GLA FACTOR VIIA	HEAVY CHAIN; CHAIN; H.	I; DES-GLA FACTOR VIIA	(LIGHT CHAIN); CHAIN: L,	M; (DPN)-PHE-ARG; CHAIN:	C, D; PEPTIDE E-76; CHAIN:	Λ, Ι,	DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H,	I, DES-GLA FACTOR VIIA	(LIGHT CHAIN); CHAIN: L,	M; (DPN)-PHE-ARG; CHAIN:	(c, D, rer 11DE E-76; ChAUN.) X, Y;	DES-GLA FACTOR VIIA	(HEAVY CHAIN); CHAIN: H,	G DES-GLA FACTOR VIIA	M. (DPN)-PHE-ARG: CHAIN. L.	C. D. PEPTIDE E-76: CHAIN:	X, Y;	DES-GLA FACTOR VIIA	(HEAVY CHAIN); CHAIN: H.	I DESTOCA FACTOR VITA
SeqFold	Score																									-				
PMF	score		0.07			0.15				0.05	5.5					10,0	-0.07					-0.15						-0.19		
Verify	score		0.1			-0.08				0.48	5					,	0.12					0.05						0.25		
PSI-	BLASI		3.60E-14			1.40E-12				3 90E_21	2.705.5						3.60E-14					3.60E-12						1.30E-13		
End	{		3976			3897				3047	1						3976					4168						4259		
Start	V		3900			3831				2858	n Cn C					1	3900					4078						4170		
Chain	2		ا ا			T					د						1					L						T		
PDB	a		ldan			ldva				Idva	,						Idva					ldva						ldva		
SEQ	a ö		808			808				800	6						608					808						808		

PDB annotation		CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECADI2, CADHERIN, CELL
Coumpound	M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN: X, Y;	E-CADHERIN; CHAIN: A, B;	E-CADHERIN, CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;				
SeqFold score									
PMF score		-	96.0	0.93	0.89	0.65	-1.41	0.25	0.99
Verify score		0.35	0.16	0.23	0.3	0.03	0.19	-0.18	0.21
PSI- BLAST		1.10E-33	9.00E-30	1.60E-20	3.60E-54	7.20E-32	3.60E-33	3.60E-28	1.80E-27
End		1234	1338	1440	1547	1652	1750	1860	1960
Start		1026	1171	1279	1352	1455	1589	1690	1780
Chain ID		ď	<	V	∢	<	∢	∢	Y
PDB ID		ledh	ledh	ledh	lcdh	ledh	ledh	ledh	ledh
SEQ D		808	608	809	808	808	608	808	608

PDB annotation	ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM
Coumpound		E-CADHERIN, CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;	E-CADHERIN, CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;				
SeqFold score									
PMF score		0.07	0.94	0.34	0.92	-	-1.41	0.86	0.75
Verify		0.17	0.3	0.13	0.19	0.08	0.37	0.07	0.14
PSI- BLAST		5.40E-20	1.80E-30	1.80E-50	9.00E-29	1.80E-38	1.80E-32	3.60E-29	1.60E-39
End		354	2062	2163	2264	2371	2473	2577	2683
Start		182	1898	1975	2104	2178	2306	2414	2488
Chain ID		<	4	∢	<	∢	4	⋖	A
PDB ID		1cdh	1edh	ledh	ledh	ledh	1edh	ledh	1edh
SEQ	Ö	808	809	809	608	808	808	808	809

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PDB annotation	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADDESIGN FROILIN, CALCIONI BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAI CADHEBIN DOMAINS	AND 2 ECADI2: CADHERIN CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	AND 2 ECADIS: CADRED IN CELL	AND 2, ECADIZ, CADRENII, CELE	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECADI2; CADHEKIN, CELL	ADHESION PROTEIN, CALCIUM BUNINNG PROTEIN	DINDING FROIEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM	BINDING PROTEIN	CELL ADHESION PROTEIN	EPITHELIAL CADHERIN DOMAINS I	AND 2, ECAD12; CADHERIN, CELL	ADHESION PROTEIN, CALCIUM BINDING PROTEIN
Coumpound		E-CADHERIN; CHAIN: A, B;			E-CADHERIN; CHAIN: A, B;					E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;					E-CADHERIN; CHAIN: A, B;				E CADITERNI CITAIN A B.	E-CADHEKIN; CHAIN: A, B;				E-CADHERIN; CHAIN: A, B;			
SeqFold score					120.62																														
PMF		0.99		_						_					1202.08			0.94					0.41				,	9.0				0.59	_		
Verify score		0.33								0.52				0.46				0.31					0.15				900	0.28				0.15			
PSI- BLAST		7.20E-31			5.40E-58					5.40E-58				1.80E-33				1.80E-28					5.40E-24				1 900	1.80E-27				1.80E-28			
End		2789			2895					2898				3003				3105					450					3191				3313			
Start AA		2619			2692					2693				2831				2941					296				3000	3045				3147			
Chain ID		4			V	;				4				٧				A					4					< _				A			
PDB ID		1edh			ledh					ledh				ledh				1cdh			_		ledh				1	ledh				ledh			
SEQ ID	5	808			808					808				809				809					608				9	608				808			

PDB annotation	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	MATRIX PROTEIN EXTRACELLULAR
Coumpound	E-CADHERIN: CHAIN: A, B;	E-CADHERIN; CHAIN: A, B,	E-CADHERIN; CHAIN: A, B;	FIBRILLIN; CHAIN: NULL;					
SeqFold score									
PMF	86.0		0.3	0.89	0.77	-	_	0.52	0.48
Verify score	0.53	0.47	0.12	0.31	0.5	0.33	0.14	0.24	90.0
PSI- BLAST	1.40E-35	1.80E-16	3.60E-53	5.40E-30	3.60E-27	1.10E-57	1.80E-33	1.80E-30	3.60E-16
End	3418	3523	248	556	662	917	1022	1129	3931
Start	3225	3355	39	401	465	718	854	959	3860
Chain ID	∢	⋖	4	<	¥	< _	4	∢	
PDB ID	1edh	ledh	ledh	ledh .	ledh	ledh	ledh	ledh	lem
SEQ NO:	809	809	608	808	808	608	608	608	808

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PDB annotation	MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN	MÁTRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT, MATRIX PROTEIN	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIGAND), BLOOD COAGULATION, 2 SERINE PROTEASE, COMPLEX, CO-FACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, EGF, COMPLEX (SERINE 4 PROTEASE/COFACTOR/LIGAND), BLOOD CLOTTING	GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13	CELL ADHESION PROTEIN CADHERIN INCG 13
Coumpound		FIBRILLIN; CHAIN: NULL;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; \$L15; CHAIN: I;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; 5L15; CHAIN: 1;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NUI.L; N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3	N-CADHERIN; INCG 3
SeqFold										
PMF		-0.18	-0.01	-0.19	-0.2	-0.2 0.63	0.87	0.3	0.72	60:0
Verify score		0.02	0.12	0.2	0.07	0.07	0.15	0.34	0	0.15
PSI- BLAST		9.00E-14	3.60E-14	1.30E-13	1.80E-17	1.80E-18 1.60E-05	5.40E-05	3.60E-17	5.40E-05	3.60E-05
End AA		4217	3976	4259	4228	4262	1232	1439	1546	229
Start		4126	3900	4170	4074	4134	1169	1350	1455	155
Chain ID			1	1						
PDB ID	c	lem n	·Ifak	1fak	lklo	1klo Incg	lncg	lncg	Incg	Incg
SEQ D NO:		608	608	608	808	809	809	608	608	809

SEO	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
8	£	9	AA	AA	BLAST	score	score	score		
.008 809	Incg		1599	1651	9.00E-06	-0.45	0.04		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	Incg		1991	1748	3.60E-06	0.35	0.31		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	Incg		1975	2061	1.80E-15	-0.06	0.1		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	lncg		2079	2161	0.00036	0.31	0.29		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	Incg		2180	2922	9.00E-07	0.11	0.01		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	lncg		2304	2369	1.80E-06	0.43	0.64		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	lncg		2411	2458	0.0045	-0.23	0.22		N-CADHERIN; 1NCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
809	lncg		2592	2681	0.00014	0.41	0.25		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
809	Incg		2692	2788	3.60E-20	0.44	0.7		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	lncg		3039	3106	9.00E-07	0.45	0.51		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	lncg		3120	3191	0.00018	90.0	0.34		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
809	Incg		3225	3312	1.10E-12	0.28	0.45		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
809	lncg		716	811	1.10E-21	0.16	8.0		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
809	lncg		852	006	0.00036	-0.19	0.53		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
608	lncg		932	1021	1.60E-05	0.55	0.46		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
809	Inci	В	1065	1129	3.60E-06	90:0	0.78		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	1178	1234	1.80E-05	-0.16	6.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
809	Inci	В	1350	1440	3.60E-16	0.53	90.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	1481	1547	5.40E-05	0.21	91.0	3	N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
809	Inci	В	1991	1750	1.40E-06	0.26	0.64		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
808	Inci	В	181	248	5.40E-05	-0.17	0.4		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN

SEQ ID NO:	PDB ID	Chain TD	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
;										INCI 13
608	Inci	В	1912	1960	0.0045	-0.5	0.27		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	1975	2062	1.80E-14	0.24	0.07		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2180	2264	3.60E-06	-0.4	0.3		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2307	2371	5.40E-07	0.39	9.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2412	2458	0.0079	-0.03	0.49		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2692	2789	1.80E-19	0.56	1		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	2832	2898	1.80E-06	-0.01	0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	æ	2951	3003	0.0045	-0.24	0.78		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	æ	3041	3105	1.10E-06	0.52	99.0		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	3146	3191	0.0013	-0.2	0.17		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	3225	3313	1.40E-10	0.31	0.17		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	716	812	5.40E-20	0.01	0.88		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	В	853	917	7.20E-05	0.26	0.84		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Inci	æ	932	1022	7.20E-06	0.4	0.59		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
608	Incj	V	1039	1234	1.10E-35	0.39	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
608	Incj	¥	1144	1338	1.40E-31	0.53	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
809	Incj	¥	1251	1440	1.80E-23	0.1	0.16		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
608	Incj	٧	1351	1547	1.60E-58	0.28	0.99		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
808	Incj	A	1455	1652	3.60E-35	0.02	0.88		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
608	Incj	٧	1562	1750	1.10E-33	0.46	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN

PDB Chain St		St	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
- AA	¥			₹	BLAST	score	score	score		
Incj A 1667 1	1991			1861	7.20E-33	0.2	0.84		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 173 3	173 3	3	3	\$	3.60E-22	80.0	0.23		N-CADHERIN, CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 1793 I	1793	-		096	9.00E-27	0.31	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 1898 2	8681		7	2062	3.60E-31	0.25	6.0		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 1975 2	1975		7	2163	1.60E-56	0.02	0.45		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2079 2	2079		12	2264	9.00E-31	0.23	0.89		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
lncj A 2180 2	2180		7	2371	3.60E-40	0.25	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2300 2	2300		117	2473	3.60E-35	0.32	-		N-CADHERIN: CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2407 2	2407		7	2577	9.00E-31	-0.03	8.0		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2488 2	2488 2	- 2	2	683	3.60E-40	90:0	9.0		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 256 4	256 4	4	4	20	1.30E-26	0.28	0.51		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2592 27	2592		27	2789	1.40E-31	0.23	-		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2691 28	2691		~	2897	3.60E-63			120.25	N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2693 28	2693		2	2898	3.60E-63	0.28	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 2825 3	2825		E .	3003	9.00E-36	0.28	0.99		N-CADHERIN, CHAIN: A,	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
	2915		` .	3105	9.00E-30	0.35			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3039	3039			3196	1.30E-30	0.16	0.8		N-CADHERIN, CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3141	3141			3313	1.30E-30	0.35	0.82		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3225	3225			3418	1.80E-39	99.0			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3349	3349			3523	1.80E-17	9.0	_		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
Incj A 3473 3	3473		[]	129	5.40E-10	0.35	-0.03		N-CADHERIN; CHAIN: A,	CELL ADHESION PROTEIN CELL

PDB annotation	ADHESION PROTEIN										7	IN; CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION		CELL ADJUGGION LIVONADILI DI
Coumpound		N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN; CHAIN: A;	N-CADHERIN, CHAIN: A,	N-CADHERIN, CHAIN: A.	N-CADHERIN; CHAIN: A;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN. L; COAGULATION FACTOR VIIA (HEAVY CHAIN); CHAIN: H; TRIPEPTIDYL INHIBITOR; CHAIN: C;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	CDITTLE IAI CADUEDINI
SeqFold score														
PMF		0.21	0.36	96.0	0.01	-	0.88		0.57	-0.08	-0.15	0.13	0.27	27.0
Verify		-0.28	80.0	0.36	-0.13	0.39	0.35	0.48	89.0	10.0	0.08	0.33	0.24	200
PSI- BLAST		1.10E-59	1.80E-35	7.20E-28	1.80E-25	1.10E-63	1.80E-34	3.60E-34	1.20E-21	3.60E-13	7.20E-11	7.20E-08	7.80E-20	1 205 17
End		248	556	799	812	917	1022	1129	3947	3976	4168	1133	1133	1230
Start		39	390	467	573	717	827	932	3865	3904	4082	1026	1041	1145
Chain		A	V	4	A	4	¥	4	٦	1	1			
PDB ID		Incj	lncj	Incj	lncj	Incj	Incj	Incj	1qfk	1qfk	lqfk	Isuh	Isuh	-
SEQ ID	NO.	608	608	608	608	809	808	809	808	808	808	809	808	000

SEQ ID	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
Ö										ADHESION
809	Isuh		1171	1238	7.20E-08	-0.03	0.4		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		1249	1334	1.30E-10	0.54	0.62		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		1350	1444	1.80E-21	0.3	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		1455	1549	1.30E-08	0.45	0.63		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	lsuh		1455	1551	5.40E-07	0.14	0.11		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		155	226	2.60E-07	0.35	0.07		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		1570	1650	1.00E-12	0.41	0.31		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1 suh		1589	1656	1.40E-08	0.2	60:0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	l suh		1667	1754	1.30E-12	0.07	0.68		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERN, CALCTUM BINDING, CELL ADHESION
809	1suh		0691	1754	3.60E-06	90.0	99:0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		1777	1867	1.30E-14	-0.06	0.48		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	Isuh		1780	1868	3.60E-05	-0.24	0.39		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		182	252	5.40E-07	-0.41	0.13		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		1898	1964	1.60E-05	-0.1	0.18		EPITHELIAL CADHERIN;	CELL ADHESION UVOMORULIN;

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ΑŞ	8	<u> </u>	Ψ¥	¥	BLAST	score	score	score		
									CHAIN: NULL;	CADHERIN, CALCIUM BINDING, CELL ADHESION
608	lsuh		5261	2066	7.20E-18	0.17	1202.08		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	lsuh		2084	2163	2.60E-15	0.42	60:0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		2104	2167	3.60E-07	0.05	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL. ADHESION
608	1 suh		2178	2268	1.30E-14	-0.23	0.57		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2178	2268	3.60E-10	-0.36	0.46		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		2280	2375	3.90E-20	0.44	0.77		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		2306	2375	1.10E-08	0.4	0.94		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UYOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	lsuh		2395	2475	3.90E-05	0.03	0.58		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2414	2477	0.00011	-0.37	90.0	-	EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		2488	2581	3.60E-12	-0.2	0.03		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	Isuh		2489	2581	1.30E-13	0.16	0.98		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	Isuh		2591	2681	2.60E-10	0.02	0.12		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2692	2793	1.40E-23	0.51	66:0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION

SEQ ID	PDB	Chain ID	Start	End	PSI- BLAST	Verify	PMF score	SeqFold	Coumpound	PDB annotation
%08 809	lsuh		2806	2902	2.60E-21	0.43	0.87		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	Isuh		2831	2902	1.40E-08	-0.01	0.53		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1 suh		2914	3007	1.20E-14	0.37	0.93		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN. CADHERIN, CALCIUM BINDING, CELL ADHESION
808	Isuh		2941	3007	1.80E-05	-0.07	0.86		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	1suh		3026	3109	2.60E-21	0.36	0.89		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1suh		3045	3109	3.60E-06	0.33	0.55		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		3120	3213	3.90E-12	0.43	0.25		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING; CELL ADHESION
808	1suh		3225	3317	1.30E-13	9.0	0.82		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
808	lsuh		3330	3422	3.90E-21	0.58	86.0		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	1suh		3355	3422	5.40E-05	9.0	0.94		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	l suh		3435	3524	6.50E-11	0.56	0.65		EPITHELIAL CADHERIN; CHAIN: NULL:	CELL ADHESION UVOMORULIN; CADHERIN. CALCIUM BINDING, CELL ADHESION
608	lsuh		363	454	0.0001	0.08	0.24		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
608	l suh		401	454	9.00E-06	-0.21	0.3		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
809	1 suh		467	260	1.30E-15	-0.03	0.19		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCTUM BINDING, CELL

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PDB annotation	ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	PLASMINOGEN ACTIVATION	BLOOD COAGULATION FACTOR STUART FACTOR; BLOOD COAGULATION FACTOR, SERINE PROTEINASE, EPIDERMAL 2 GROWTH FACTOR LIKE DOMAIN		CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN			
Coumpound		EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	EPITHELIAL CADHERIN; CHAIN: NULL;	T-PLASMINOGEN ACTIVATOR FI-G; ITPG 7 CHAIN: NULL; ITPG 8	BLOOD COAĞULATION FACTOR XA; CHAIN: L, C;	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	E-CADHERIN; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;
SeqFold												
PMF		0.57	0.95	0.72	0.47	0.57	0.18	0.59	0.1	-0.2	0.25	0.16
Verify score		0.39	-0.21	0.4	0.27	0.37	0.2	0.21	-0.26	0.02	-0.26	0.35
PSI- BLAST		3.90E-05	7.20E-25	1.00E-17	1.30E-07	5.20E-13	9.00E-07	1.30E-18	7.20E-11	5.40E-12	3.60E-24	1.80E-17
End		859	816	921	921	1026	1026	3940	3994	4234	370	145
Start AA		965	716	826	854	930	656	3863	3904	4069	190	23
Chain									L	A	4	¥
PDB		Isuh	Isuh	1 suh	1suh	lsuh	1 suh	Itpg	lxka	9wga	ledh	ledh
SEQ ID	:0N	809	808	809	809	608	809	808	808	808	811	811

1	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
	1cdh	<	296	474	1.80E-30	0.21	0.82		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
	ledh	«	410	584	7.20E-26	0.21	-		E-CADHERIN, CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
1	1edh	<	50	237	1.80E-51	0.61	_		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
+	ledh	<	50	254	1.80E-51			124.11	E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS I AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
+-	Incg		188	238	0.00018	0.07	0.03		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
+	Incg		408	473	1.80E-06	0.34	0.63		N-CADHERIN, INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
+	lncg		48	143	1.60E-19	0.16	-		N-CADHERIN; INCG 3	CELL ADHESION PROTEIN CADHERIN INCG 13
+	Inci	В	189	238	0.00036	-0.15	0.12		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
+	Inci	В	409	474	3.60E-07	0.44	0.62		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
+	Inci	В	48	145	5.40E-19	0.35	-		N-CADHERIN; INCI 3	CELL ADHESION PROTEIN CADHERIN INCI 13
+	lncj	<_	160	370	1.10E-26	0.21	0.98		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
+	Incj	₹	24	145	3.60E-20	0.31	0.59		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
+	Incj	<	27.1	474	1.80E-31	0.31	0.58		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
	1ncj	<	402	999	7.20E-28	0.24	66.0		N-CADHERIN: CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
- 1	- Ju	<	49	238	1.80E-55	0.29			N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL

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PDB annotation	ADHESION PROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION ÜVOMORULIN, CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN: CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound		N-CADHERIN; CHAIN: A;	EPITHELIAL CADHERIN; CHAIN: NULL;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	QGSR ZINC FINGER PEPTIDE: CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B. D, E. CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F. G;							
SeqFold score		125.8											
PMF score			0.18	0.23	0.01	0.46	68.0	_	_	0.1	-	0.92	-
Verify score			0.22	-0.08	0.36	0.3	-0.15	0.65	0.58	0.23	0.61	0.68	0.34
PSI- BLAST		1.80E-55	1.20E-16	0.0041	5.20E-12	1.30E-17	1.60E-08	9.00E-23	2.60E-29	1.00E-08	2.60E-35	3.60E-27	7.20E-49
End AA		253	258	237	371	478	478	149	149	584	142	142	861
Start AA		49	162	190	268	383	410	48	49	490	62	69	117
Chain ID		¥									A	Y	ပ
PDB ED		lncj	Isuh	1suh	l suh	1suh	Isuh	1suh	1suh	1suh	1a1h	lalh	Ime y
SEQ NO:		811	118	811	118	811	811	118	811	811	813	813	813

PDB annotation		ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III. 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	DOMAIN) MUTANT WITH IPAA 4 PRO 131 REPLACED BY ALA, PRO 133 REPLACED BY ALA, CYS 140 IPAA 5 REPLACED BY ALA (P131A,P133A,C140A) (NMR, 10 STRUCTURES) IPAA 6	SP1F2; CHAIN: NULL;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score				111.05			
PMF		0.35	0.98		_	-	
Verify		0.27	0.17		0.36	0.38	0.26
PSI- BLAST		5.40E-07	1.80E-38	2.60E-61	9.00E-36	7.20E-37	2.60E-49
End		312	270	310	312	235	254
Start		286	118	145	174	06	144
Chain ID			<	<	∢	<	O
PDB ID		1sp2	1116	1116	1116	1116	lubd
SEQ ID NO:		813	813	813	813	813	813

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIM: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIM: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score		105.14				
PMF	_		1	-	0.94	0.98
Verify	0.25		0.03	0.37	0.49	0.34
PSI- BLAST	3.60E-35	1.30E-50	7.80E-50	1.30E-50	2.60E-43	1.80E-33
End	254	283	283	310	170	170
Start	153	172	178	661	99	69
Chain ID	O	ပ	U	ပ	U	U
PDB ID	lubd	lubd	lubd	lubd	lubd	lubd
SEQ ID NO:	813	813	813	813	813	813

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
e ë	e	<u>e</u>	*	¥¥	BLAST	score	score	score		
										(TRANSCRIPTION REGULATION/DNA)
813	1ubd	ပ	97	861	1.80E-34	0.62			YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION
									ASSOCIATED VIRUS PS	KEGULATION/DNA) YING-YANG I;
									CHAIN: A B:	IKANSCKIPTION INTERATION,
									CIONIC O, D,	FINGER PROTEIN DNA-PROTEIN
										RECOGNITION, 3 COMPLEX
										(TRANSCRIPTION REGULATION/DNA)
813	2adr		62	116	2.60E-23	0.74	_		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION
										TRANSCRIPTION REGULATION,
										ADRI, ZINC FINGER, NMR
813	2gli	¥	145	284	3.90E-62			107.07	ZINC FINGER PROTEIN GLII;	COMPLEX (DNA-BINDING
									CHAIN: A; DNA; CHAIN: C,	PROTEIN/DNA) FIVE-FINGER GLI; GLI,
								_	D;	ZINC FINGER, COMPLEX (DNA-
										BINDING PROTEIN/DNA)
813	2gli	<	145	312	3.90E-62	0.23	0.98		ZINC FINGER PROTEIN GLII;	COMPLEX (DNA-BINDING
									CHAIN: A; DNA; CHAIN: C,	PROTEIN/DNA) FIVE-FINGER GLI; GLI,
									.; ```	ZINC FINGER, COMPLEX (DNA-
				1						BINDING PROTEIN/UNA)
813	2gli	٧_	<u></u>	309	5.40E-34	0.39			ZINC FINGER PROTEIN GLII;	COMPLEX (DNA-BINDING
									CHAIN: A; DNA; CHAIN: C,	PROTEIN/DNA) FIVE-FINGER GLI, GLI,
									Ω	ZINC FINGER, COMPLEX (DNA-
			ļ	١						BINDING PROJEINUNA)
<u> </u>	Zgli	<_	62	172	3.90E-43	0.41	_		ZINC FINGER PROTEIN GLII,	COMPLEX (DNA-BINDING
									CHAIN: A; DNA; CHAIN: C,	PROTEIN/DNA) FIVE-FINGER GLI, GLI,
									ä	ZINC FINGER, COMPLEX (DNA-
013	12.0		33		יייייייייייייייייייייייייייייייייייייי				The second and second of the	BINDING PROTEIN/UNA)
610	11.87	<	?	109	1.305-30	0.47	-		CHAPIT A: DNA: CHAPIT.	COMPLEX (DNA-BINDING
			_						CIPAIN: A, DIVA, CIPAIN: C,	ZINCI EINCER COMPLEX (DNA.
									î	BINDING PROTEIN/DNA)
813	2gli	٧	68	256	9.10E-60	0.4	_		ZINC FINGER PROTEIN GLII;	COMPLEX (DNA-BINDING
									CHAIN: A; DNA; CHAIN: C,	PROTEIN/DNA) FIVE-FINGER GLI; GLI,
									Ď.	ZINC FINGER, COMPLEX (DNA-
			-							BINDING PROTEIN/DNA)
813	2gli	⋖	97	225	1.80E-33	0.45	0.94		ZINC FINGER PROTEIN GLII;	COMPLEX (DNA-BINDING
									CHAIN: A; DNA; CHAIN: C,	PROTEIN/DNA) FIVE-FINGER GLI; GLI,
										ZINC FINGER, COMPLEX (DNA-
814	laka	-	,	386	0.005 50	20.0	0.34		LINDAID A DODINING NI	TO ANOCEED A OF CAMA DINIDING
	1800	-	7	200	7.00E-30	-0.02	0.04		HINKING ARGINING IN-	I KANSFEKASE SAM-BINDING

PDB annotation	E; DOMAIN, BETA-BARREL, MIXED ALPHA-BETA, HEXAMER, 2 DIMER	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT				GOMPLEX (GTP- BINDING/TRANSDUCER) BETA I, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION		Ö	N: ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	NGE COMPLEX (HSP24/HSP70) HSP70,
Coumpound	METHYLTRANSFERASE; CHAIN: 1, 2, 3, 4, 5, 6;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUPI; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA: CHAIN: G;	KARYOPHERIN BETA2; CHAIN: B; RAN; CHAIN: C;	BETA-CATENIN; CHAIN NULL;	NUCLEOTIDE EXCHANGE FACTOR GRPE: CHAIN: A. B.
SeqFold score										78.03
PMF		0.98	0.03	-	_	ļ	-	0.07	0.23	
Verify score		0.32	0.16	0.12	0.28	0.18	0.63	-0.04	-0.02	
PSI- BLAST		1.20E-07	0.0036	3.60E-60	3.60E-68	3.60E-60	3.60E-73	5.20E-05	2.60E-15	5.20E-41
End		520	1078	1099	1147	9601	1144	653	653	222
Start AA		235	821	196	831	758	823	861	345	51
Chain ID		₹	<	<	¥	В	a	В		∢
PDB ID		1b3u	lcrz	lerj	lcrj	lgot	lgot	1qbk	3bct	1dkg
SEQ ID	Ë	817	817	817	817	817	817	817	817	818

SEQ	PDB	Chain ID	Start	End	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
Ö									DNAK; CHAIN: D;	COILED-COIL, COMPLEX (HSP24/HSP70)
818	1dkg	<	53	221	1.80E-20	-0.15	0.83		NUCLEOTIDE EXCHANGE FACTOR GRPE, CHAIN: A, B; MOLECULAR CHAPERONE DNAK; CHAIN: D;	COMPLEX (HSP24/HSP70) HSP70, GRPE, MOLECULAR CHAPERONE, NUCLEOTIDE EXCHANGE 2 FACTOR, COILED-COIL, COMPLEX (HSP24/HSP70)
818	ldkg	4	19	220	5.20E-41	-0.29	0.42		NUCLEOTIDE EXCHANGE FACTOR GRPE: CHAIN: A. B; MOLECULAR CHAPERONE DNAK: CHAIN: D;	COMPLEX (HSP24/HSP70) HSP70, GRPE, MOLECULAR CHAPERONE, NUCLEOTIDE EXCHANGE 2 FACTOR, COILED-COIL, COMPLEX (HSP24/HSP70)
										Ciddiga tiral yarras co
820	lan2	¥	135	203	3.60E-14	-0.73	0.03		MAX PROTEIN; CHAIN: A, C; DNA; CHAIN: B, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) MYN PROTEIN; MAX, DNA BINDING, BASIC-HELIX-LOOP- HELIX-LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)
820	lan2	K	140	219	2.60E-13	8.0-	60:00		MAX PROTEIN; CHAIN: A, C; DNA; CHAIN: B, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) MYN PROTEIN; MAX, DNA BINDING, BASIC-HELIX-LOOP- HELIX-LEUCINE ZIPPER, 2 TRANSCRIPTION FACTOR, COMPLEX (DNA-BINDING PROTEIN/DNA)
820	1hlo	4	131	203	1.60E-14	-0.58	0		TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5'- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	COMPLEX (TRANSCRIPTION FACTOR MAX/DNA) TRANSCRIPTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAX/DNA)
820	1hlo	В	132	203	1.80E-14	-0.27	0.28		TRANSCRIPTION FACTOR MAX; CHAIN: A, B; DNA (5'- D(*CP*AP*CP*CP*AP*CP*GP *TP*GP*GP*T)-3', CHAIN: C, D;	COMPLEX (TRANSCRIPTION FACTOR MAX/DNA) TRANSCRIPTIONAL REGULATION, DNA BINDING, COMPLEX 2 (TRANSCRIPTION FACTOR MAX/DNA)
	_									
824	1b3u	<	44	373	6.50E-08	0.55	_		PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT
824	1 ibr	В	131	235	0.0026	0.25	0.82		RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT;	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR

SEQ ID	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									CHAIN: B, D;	TRANSPORT RECEPTOR
824	1ibr	В	201	303	6.50E-06	90.0	0.47		RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR
824	3bct		154	404	6.50E-14	0.41	66.0		BETA-CATENIN; CHAIN: NULL;	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
826	168g	<	70	294	5.40E-14	0.41	0.22		I-AMINOCYCLOPROPANE-I- CARBOXYLATE SYNTHASE; CHAIN: A, B;	LYASE ACC SYNTHASE, S- ADENOSYL-L-METHIONINE ETHYLENE BIOSYNTHESIS
826	1691	<	76	317	3.60E-37	0.12	-0.13		3-AMINO-5- HYDROXYBENZOIC ACID SYNTHASE; CHAIN: A;	RIFAMYCIN BIOSYNTHESIS (RIFD GENE) AHBA SYNTHASE; RIFAMYCIN BIOSYNTHESIS (RIFD GENE)
826	1bj4	4	_	312	5.40E-50	0.25	0		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A;	TRANSFERASE, METABOLIC ROLE, PYRIDOXAL 5'-PHOSPHATE
826	1bjw	<	8	309	1.80E-52	0.29	1202.08		ASPARTATE AMINOTRANSFERASE; CHAIN: A, B;	AMINOTRANSFERASE AMINOTRANSFERASE, PYRIDOXAL ENZYME
826	1680	∢		317	7.20E-06		·	52.33	8-AMINO-7-OXONANOATE SYNTHASE, CHAIN: A;	TRANSFERASE AONS, 8-AMINO-7- KETOPELARGONATE SYNTHASE; PLP-DEPENDENT ACYL-COA SYNTHASE, BIOTIN BIOSYNTHESIS, 8-2 AMINO-7-OXONANOATE SYNTHASE, 8-AMINO-7- KETOPELARGONATE 3 SYNTHASE, TRANSFERASE
826	1c0n	Ą	10	311	9.00E-57	0.51	_		CSDB PROTEIN; CHAIN: A;	LYASE ALPHA/BETA FOLD
826	1cj0	¥	-	312	3.60E-51	0.04	-1.41		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B;	TRANSFERASE SHMT; HYDROXYMETHYL TRANSFERASE, I CARBON METABOLISM
826	101	¥	93	314	3.60E-16	0.1	-0.19	·	CYSTATHIONINE BETA- LYASE: CHAIN: A, B;	METHIONINE BIOSYNTHESIS BETA CYSTATHIONASE, PLP-DEPENDENT ENZYMES, METHIONINE BIOSYNTHESIS, C-S BETA 2 LYASE
826	1cs1	4	22	313	5.40E-34	0.2	-0.03		CYSTATHIONINE GAMMA- SYNTHASE; CHAIN: A, B, C, D;	LYASE CGS; LYASE, LLP-DEPENDENT ENZYMES, METHIONINE BIOSYNTHESIS
826	1d2f	<	135	259	1.80E-09	-0.06	0.51		MALY PROTEIN; CHAIN: A, B;	TRANSFERASE AMINOTRANSFERASE FOLD, LARGE PLP-BINDING DOMAIN,

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ΕŞ	8	£	¥	ΥΥ	BLAST	score	score	score		
										SMALL C- 2 TERMINAL DOMAIN, OPEN ALPHA-BETA STRUCTURE.
826	1dfo	<	∞	314	9.00E-54	-0.07	0.17		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D;	TRANSFERASE SHMT, SERINE METHYLASE; ALPHA PLP ASPARTATE, AMINO TRANSFERASE, (AAT)-LIKE FOLD
826	leg5	4	32	317	3.60E-58	0.62	-		AMINOTRANSFERASE; CHAIN: A, B;	TRANSFERASE PLP-DEPENDENT ENZYMES, IRON-SULFUR-CLUSTER SYNTHESIS, C-S 2 BETA LYASE
826	leji	≺	-	312	5.40E-50	0.09	-1.41		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D;	TRANSFERASE SHMT; SERINE- GLYCINE CONVERSION, PYRIDOXAL 5-PHOSPHATE, 2 TETRAHYDROFOLATE, ASYMMETRIC DIMER
826	lqgn	¥.	20	311	7.20E-21	0.19	-0.06		CYSTATHIONINE GAMMA- SYNTHASE; CHAIN: A, B, C, D, E, F, G, H;	LYASE METHIONINE BIOSYNTHESIS, PYRIDOXAL 5'-PHOSPHATE, GAMMA- 2 FAMILY, LYASE
826	1tpl	<	101	311	3.60E-09	0.21	-0.03		LYASE(CARBON-CARBON) TYROSINE PHENOL-LYASE (E.C.4.1.99.2) 1TPL 3	
826	2tpl	4	101	311	1.30E-08	0.02	-0.06		TYROSINE PHENOL-LYASE; CHAIN: A, B;	LYASE LYASE, PLP-DEPENDENT ENZYME, PYRIDOXAL PHOSPHATE
830	1c4z	¥.	1213	1566	1.10E-45			217.98	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
830	1c4z	⋖	1239	1565	1.10E-45	0.38			UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP, UBCH7, BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
830	1c0	<	608	841	2.60E-11	90:0	0.29		WWPROTOTYPE; CHAIN: A;	SH3 PROTOTYPE WWPROTOTYPE, PROTEIN DESIGN
830	п В	4	586	1019	3.90E-12	0.01	0.35		WWPROTOTYPE; CHAIN: A;	SH3 PROTOTYPE WWPROTOTYPE, PROTEIN DESIGN
832	lalh	A	179	259	3.60E-26	-0.37	0.24		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
832	laih	A	235	315	2.60E-39	0.04	_		QGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
S S	1	3	{	ŧ	TCV10	3000	31036	3006		
									PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
832	lath	<	263	344	9.10E-38	0.48	_		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
832	lalh	∢	598	678	1.40E-30	0.41	66.0			COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER. DNA-BINDING PROTEIN
832	lalh	Y	598	678	3.90E-39	0.27	0.98		QGSR ZINC FINGER PEPTIDE: CHAIN: A: DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
832	Ime y	U	150	231	3.60E-40	-0.35	0.07		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
832	lme y	U	178	259	1.10E-43	-0.16	0.52		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
832	Ime y	U	206	287	3.60E-45	0.1	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
832	Ime y	ပ	234	315	1.60E-46	0.23	~		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
832	Ime y	ပ	262	343	1.40E-47	0.55	-		DNA: CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER. PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
832	Imc	C	290	371	1.30E-48	0.49	-		DNA; CHAIN: A, B, D, E,	COMPLEX (ZINC FINGER/DNA) ZINC

Chain ID		Start	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound CONSENSUS ZINC FINGER	PDB annotation FINGER, PROTEIN-DNA
								PROTEIN; CHAIN: C, F, G;	NTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 318	318		399	1.80E-49	0.62	_		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
346	346		427	1.40E-49	0.34	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 374	374		455	3.60E-50	0.27	_		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 402	402		483	5.40E-50	0.42	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 430	430		511	7.20E-50	0.16	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 430	430		538	6.50E-33	-0.22	0.98		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 458	458		539	1.10E-49	0.1	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C 514	514		594	7.20E-48	0.43	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA

PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIALION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	ON THE PROPERTY OF THE PROPERT	COMPLEX (IKANSCRIPTION	REGULATION/DNA) COMPLEX	(I'KANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POI YMERASE III 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA
Coumpound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE:	CHAIN: B, C, E, F,				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE,	CHAIN: B, C, E, F;			0. 4 1 14110 1110	IFIIIA; CHAIN: A, D, 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B. C. E. F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;	
SeqFold score							105.86		•	-	-		-																											
PMF		0.84											_						-1.41						-1.41						68.0						98.0			
Verify		60.0											CI.0						0.32						0.16						-0.05						-0.04			
PSI- BLAST		1.80E-36					1.30E-66					75 707 .	1.60E-36						3.60E-37						1.30E-36						1.60E-36						1.80E-35			
End		352					431					90,	408						464						575						603						631			
Start AA		207					262					.,,	707						319						431						459						487			
Chain TD		∢		_			٧						∀						٧						٧						V						<			
PDB ID		91116					1116					207.	911						1tf6						1166						9111									
SEQ ID NO:		832					832					,,,	832						832						832						832						832	_		

PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY 1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
Coumpound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A. B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score							
PMF		_	0.18	96.0	69.0	96.0	-
Verify		0.3	-0.36	-0.38	-0.37	-0 07	-0.01
PSI- BLAST		7.20E-35	9.00E-29	3.60E-31	1.30E-34	2.60E-48	3.60E-32
End		099	259	287	315	371	343
Start		515	158	186	188	232	242
Chain 1D		₹	ပ	U	U	U	ပ
PDB ID		1166	lubd	1ubd	lubd	lubd	lubd
SEQ D		832	832	832	832	832	832

	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	 e	<u>e</u>	¥ ¥	¥	BLAST	score	score	score		
-										RECOGNITION, 3 COMPLEX (TRANSCRIPTION/DNA)
ļ <u> </u>	lubd	U	270	371	1.80E-34	0.23	-		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA:	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,
									CHAIN: A, B;	INITIATOR ELEMENT, YYI, ZINC 2
										RECOGNITION, 3 COMPLEX
上	1 ubd	C	298	399	5.40E-34	0.27	-		YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION
									ASSOCIATED VIRUS PS	REGULATION/DNA) YING-YANG 1;
									CHAIN: A, B;	INITIATOR ELEMENT, YYI, ZINC 2
										FINGER PROTEIN, DNA-PROTEIN RECOGNITION. 3 COMPLEX
										(TRANSCRIPTION REGULATION/DNA)
-) pqn1	O	344	483	6.50E-52	0.04	0.88		YYI; CHAIN: C; ADENO- ASSOCIATED VIRIS P5	COMPLEX (TRANSCRIPTION REGILI ATION/DNA) VING-YANG I
					-				INITIATOR ELEMENT DNA;	TRANSCRIPTION INITIATION,
					-				CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2
										FINGER PROTEIN, DNA-PROTEIN
										(TRANSCRIPTION REGULATION/DNA)
_) pqn(C	354	455	3.60E-34	0.2	_		YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION
									ASSOCIATED VIRUS PS	REGULATION/DNA) YING-YANG 1;
									INITIATOR ELEMENT DNA;	IKANSCRIPTION INITIATION,
									,	FINGER PROTEIN, DNA-PROTEIN
										RECOGNITION, 3 COMPLEX
4	+									(TRANSCRIPTION REGULATION/DNA)
_) pqn(ပ	382	483	7.20E-36	-0.09	0.99		YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION
									ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA:	REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION.
						-			CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2
					-					FINGER PROTEIN, DNA-PROTEIN
					-					RECOGNITION, 3 COMPLEX
1-	1hd	C	410	110	1 40E 34	70.0	-		WWI. CHABI C. APPRIC	(IRANSCRIPTION REGULATION/DNA)
		-	2	-	1:40E-34	90.7	-		ASSOCIATED VIRUS P5	REGULATION/DNA) YING-YANG I:
									INITIATOR ELEMENT DNA;	TRANSCRIPTION INITIATION,
4									CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2

PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1, TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN - RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I;
Coumpound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A. B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5
SeqFold score							86.51
PMF score		0.53	0.34	_	-	0.98	
Verify		-0.42	0.02	0.23	0.22	0.43	
PSI- BLAST		3.90E-42	3.60E-32	1.30E-47	5.40E-36	5.20E-51	5.20E-51
End		594	566	622	622	678	619
Start		428	466	519	522	567	895
Chain ID		U	ပ	ပ	ပ	ပ	S
PDB ID		lubd	lubd	lubd	lubd	lubd	Jubd
SEQ ID	Ö	832	832	832	832	832	832

<u>~</u> ~	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									CHAIN: A, B;	INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
ļ	Iubd	O	577	678	5.40E-34	-0.06	_		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
I-	pqn1	U	651	089	2.60E-06	-0.04	0.18		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
1	1zfd		653	089	0.00013	0.15	0.72		SWIS; CHAIN: NULL;	ZINC FINGER DNA BINDING DOMAIN DNA BINDING MOTIF, ZINC FINGER DNA BINDING DOMAIN
1, ,	2gli	4	133	317	1.30E-36	-0.16	0.46		ZINČ FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
, ,	2gli	∢	150	286	1.60E-31	-0.24	0.49		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
1	2gli	⋖	208	345	2.60E-52	-0.21	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
	2gli	∢	235	401	5.20E-63	-0.1	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
, ,	2gli	<	262	429	3.90E-64	0.06	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)

PDB annotation		-		ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-D; D; BINDING PROTEINDNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-B); D; BINDING PROTEIN/DNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA: CHAIN: C, ZINC FINGER, COMPLEX (DNA-BINDING PROTEINDNA)	ZINC FINGER PROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: A; DNA; CHAIN: C, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	HALOALKANE HYDROLASE HYDROLASE, HALOALKANE HALOALKANE HALOALKANE DEHALOGENASE
1 C	score sco		9	6	66	61	0.19	66.0		91.33	0.12
-	score so	0.01	0.25 0.99	-0.4 0.19	0.03	-0.21 0.49	-0.07 0.	-0.02	0.53		-0.1
ŀ	PSI- BLAST	3.60E-33	2.60E-64	3.90E-57	3.60E-34	1.10E-32	5.20E-60	5.40E-31	3.90E-64	3.90E-64	0.00078
-	AA	398	485	596	482	565	652	624	678	089	1332
	Start	270	318	346	354	438	459	494	541	541	1240
	Chain ID	V	<	₹.	4	¥	V	V	4	<	
	E CI	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	2gli	166g
	SEQ ID	832 832	832	832	832	832	832	832	832	832	834

Chain Start ID AA		End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
	-						DEHALOGENASE; 1- CHLOROHEXANE CHAIN: A;	DEHALOGENASE I-S BOND
1251 13	13	362	0.00026	0.04	69.0		TRIACYLGLYCEROL HYDROLASE, CHAIN: NULL;	HYDROLASE TRIACYLGLYCEROL- HYDROLASE, X-RAY CRYSTALLOGRAPHY, 2 PSEUDOMONADACEAE, OXYANION, CIS-PEPTIDE, HYDROLASE
1251 13	23	1362	0.00026	-0.1	0.53		PALMITOYL PROTEIN THIOESTERASE 1; CHAIN: A;	HYDROLASE ALPHA/BETA HYDROLASE, GLYCOPROTEIN
1251 13	13	1362	0.00026	0.13	0.46		LACTONIZING LIPASE; CHAIN: A;	HYDROLASE TRIACYL-GLYCEROL LIPASE; LIPASE, ALPHA-BETA HYDROLASE FOLD, PSEUDOMONAS, PHOSPHONATE 2 INHIBITOR
13 241	24	_	5.40E-07	0.23	0.06		TRYPTOPHANASE; CHAIN: A, B, C, D;	TRYPTOPHAN BIOSYNTHESIS TRYPTOPHAN INDOLE-LYASE; TRYPTOPHAN BIOSYNTHESIS, TRYPTOPHAN BIOSYNTHESIS, TRYPTOPHAN INDOLE-LYASE, PYRIDOXAL 2 5-PHOSPHATE, MONOVALENT CATION BINDING SITE
13 246	246	1	7.20E-36	80.0	-0.07		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A;	TRANSFERASE TRANSFERASE, METABOLIC ROLE, PYRLDOXAL 5'- PHOSPHATE
2 244	24,	4	9.00E-41	80.0	-1.41		CSDB PROTEIN; CHAIN: A;	LYASE ALPHA/BETA FOLD
12 246	24	ب ا	7.20E-37	0.05	-0.13		SERNE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B;	TRANSFERASE SHMT; HYDROXYMETHYL TRANSFERASE, I CARBON METABOLISM
4 242	74	2	1.60E-31	-0.28	0		CYSTATHIONINE GAMMA- SYNTHASE; CHAIN: A, B, C, D;	LYASE CGS; LYASE, LLP-DEPENDENT ENZYMES, METHIONINE BIOSYNTHESIS
4 252	25	2	3.60E-40	0.17	1.41		SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D:	TRANSFERASE SHMT, SERINE METHYLASE; ALPHA PLP ASPARTATE, AMINO TRANSFERASE, (AAT)-LIKE FOLD
-	2	248	1.30E-50	0.22	_		AMINOTRANSFERASE; CHAIN: A. B:	TRANSFERASE PLP-DEPENDENT ENZYMES, IRON-SULFUR-CLUSTER SYNTHESIS, C-S 2 BETA LYASE
1 24	24	248	1.80E-45	0.1			AMINOTRANSFERASE; CHAIN: A, B;	TRANSFERASE PLP-DEPENDENT ENZYMES, IRON-SULFUR-CLUSTER SYNTHESIS, C-S 2 BETA LYASE

								<u> </u>		\neg
PDB annotation	TRANSFERASE SHMT; SERINE- GLYCINE CONVERSION, PYRIDOXAL 5-PHOSPHATE, 2 TETRAHYDROFOLATE, ASYMMETRIC DIMER	LYASE FES CLUSTER BIOSYNTHESIS, PYRIDOXAL S'-PHOSPHATE, 2 THIOCYSTEINE, AMINOACR YLATE, ENZYME-PRODUCT COMPLEX	CARBOXYLIC ESTER HYDROLASE PHOSPHOLIPASE, TRIMER, CALCIUM BINDING, ACTIVATOR SITE, 2 CARBOXYLIC ESTER HYDROLASE	HYDROLASE PLA2, PHOSPHATIDE SN-2 ACYLHYDROLASE; HYDROLASE, PHOSPHOLIPASE A2, LIPID DEGRADATION, PRESYNAPTIC 2 NEUROTOXIN, VENOM	HYDROLASE HYDROLASE, PHOSPHOLIPASE A2, PLATELET AGGREGATION INHIBITOR, 2 PBPB HEADER MODRES					LIPID DEGRADATION PHOSPHOLIPASE A2, LIPID
Coumpound	SERINE HYDROXYMETHYLTRANSF ERASE; CHAIN: A, B, C, D;	L-CYSTEINE/L-CYSTINE C-S LYASE; CHAIN: A. B;	PHOSPHOLIPASE A2; CHAIN: NULL;	PHOSPHOLIPASE A2; CHAIN: NULL;	PHOSPHOLIPASE A2; CHAIN: NULL;	HYDROLASE PHOSPHOLIPASE A2 (E.C.3.1.1.4) IPOA 3	HYDROLASE PHOSPHOLIPASE A2 (E.C.3.1.1.4) COMPLEX WITH THE 1POC 3 TRANSITION- STATE ANALOGUE 1POC 4	HYDROLASE PHOSPHOLIPASE A2 (E.C.3.1.1.4) COMPLEX WITH THE 1POC 3 TRANSITION- STATE ANALOGUE 1POC 4	HYDROLASE CALCIUM- FREE PHOSPHOLIPASE A=2= (E.C.3.1.1.4) IPP2 4	PHOSPHÖLIPASE A2; CHAIN: A, B;
SeqFold score							99.59			
PMF score	-0.11	0.83	-0.08	-0.18	-0.13	-0.14		0.21	-0.14	-0.17
Verify score	0.41	-0.03	0.38	0.07	0.07	0.3		0.08	0.2	0.12
PSI- BLAST	7.20E-35	3.90E-28	4.80E-39	4.80E-41	1.60E-40	1.60E-38	1.40E-36	0.0054	1.60E-38	3.20E-44
End	246	241	234	244	253	234	289	438	253	253
Start AA	12	-	131	131	131	131	151	355	131	131
Chain ID	∢	4							R	A
PDB ID	leji	1elu	1 a 3d	lac7	1bk9	1poa	Ipoc	lpoc	1pp2	lvap
SEQ ID	836	836	837	837	837	837	837	837	837	837

								Źω	(1)	w	TANT	TANT	DHR;	
uo	LASE	SE,	HYDROLASE HYDROLASE, LIPID DEGRADATION, CALCIUM, PRESYNAPTIC 2 NEUROTOXIN, VENOM					OXIDOREDUCTASE PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASE	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN	KINASE HCASK, GLGF REPEAT, DHR; PDZ DOMAIN, NEUREXIN, SYNDECAN, RECEPTOR CLUSTERING,	SIGNAL TRANSDUCTION HDLG
PDB annotation	I, HYDRO	IYDROLA SE A2, ANT	IYDROLA 4, CALCII 2 NEURC					TASE PD OXIDE S	OGNITION I, PROTEI	OGNITION V, PROTE N	F, CYTO S CHEMC DOMAIN	E CYTO E CHEMC DOMAIN	K, GLGF NEURE) ECEPTO	SDUCTION
PD	DEGRADATION, HYDROLASE	HYDROLASE HYDROLASE, PHOSPHOLIPASE A2, ANTICOAGULANT	HYDROLASE HYDROLASE, LIPI DEGRADATION, CALCIUM, PRESYNAPTIC 2 NEUROTOXIN, VENOM					OREDUC S, NITRIC	PEPTIDE RECOGNITION I RECOGNITION, PROTEIN LOCALIZATION	PEPTIDE RECOGNITION I RECOGNITION, PROTEIN LOCALIZATION	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTF FACTOR, PDZ DOMAIN	CYTOKINE LCF; CYTOKINE LYMPHOCYTE CHEMOATTF FACTOR, PDZ DOMAIN	KINASE HCASK, GLGF REI PDZ DOMAIN, NEUREXIN, SYNDECAN, RECEPTOR C	SIGNAL TRANSDUCTION DHR 3 DOMAIN: SIGNAL
	DEGR	HYDR PHOSI ANTI(HYDROI DEGRAI PRESYN VENOM					OXIN SON SON SON SON SON SON SON SON SON SO	RECC LOCA	RECC LOCA	CYTC LYM FACI	CYTC LYM FACT	KINA PDZ SYNI	SIGNAL
		CHAIN:	CHAIN:	XYL ASE -2- H ASP	R, SER Y, 3P2P N 67	1929 6 (62-66), 1929 7		OXIDE A: NIN: B;	RIPT;	RIPT;	HAIN:	HAIN:	Ž.	SE II I ·
Coumpound		ASE A2;	ASE A2;	E(CARBC SPHOLIP PHATIDE (OLASE)	D BY SE	BY TYR 3.1 1.4)		NITRIC CHAIN: IDE; CHA	AIN: A; C	AIN: A; C	UN 16; CI	CIN 16; CI	-2 PROTE	SCS LAR
Cou		PHOSPHOLIPASE A2; CHAIN: NULL;	PHOSPHOLIPASE A2; CHAIN: A, B;	HYDROLASE(CARBOXYL ESTER) PHOSPHOLIPASE A=2= (PHOSPHATIDE-2- ACYL-HYDROLASE) MITANT 3P2P 4 WITH ASP	59 REPLACED BY SER, SER 60 REPLACED BY GLY, 3P2P 5 63-66 DFI FTED ASN 67	REPLACED BY TYR 3P2P 6 (ID59S\$, /S60G\$, DEL(62-66), N67Y\$) (E.C.3.11.4) 3P2P 7		NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	PSD-95; CHAÎN: A; CRIPT; CHAÎN: B;	PSD-95; CHAIN: A; CRIPT; CHAIN: B;	INTERLEUKIN 16; CHAIN: NULL;	INTERLEUKIN 16; CHAIN: NULL;	HCASK/LIN-2 PROTEIN; CHAIN: A, B;	HUMAN DISCS LARGE
-		PHOSP NULL;	PHOS A, B;	HYI EST A=2 ACY	59 F 60 F 60 S	S S S		E SY E	CH	PSI	Z Z	E S	원명	田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田田
SeqFold score														
PMF score		-0.11	-0.08	-0.06				86.0	0.72	0.1	66.0	0.45	99.0	0.88
Verify		0.15	0.4	0.34				0.79	0.21	-0.25	0.59	-0.03	0.04	0.45
PSI- BLAST		4.80E-41	1.30E-40	1.60E-38				1.30E-13	1.10E-18	6.40E-10	7.20E-16	1.40E-14	1.30E-14	6.40E-17
End AA E	-	 		236 1.0			-	106 1.	102 1.	317 6.	109 7.	108	108	113 6
-	+	248	244	2			-		-					
Start		131	131	132			<u> </u>	21	12	254	21	4	21	20
Chain ID			<	¥				<	⋖	4			<	
PDB ID		lvip	2not	Зр2р				1b8q	1be9	1be9	1:16	1116	lkwa	1 pdr
SEQ	ë	837	837	837				838	838	838	838	838	838	838

					 r	_						$\overline{}$
PDB annotation	TRANSDUCTION, SH3 DOMAIN, REPEAT	OXIDOREDUCTASE BETA-FINGER	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING	HYDROLASE PDZ DOMAIN, HUMAN PHOSPHATASE, HPTP1E, PTP-BAS, SPECIFICITY 2 OF BINDING		OXIDOREDUCTASE (D, L) STEREOSPECIFIC OPINE DEHYDROGENASE, OXIDOREDUCTASE	OXIDOREDUCTASE OXYDOREDUCTASE	TRANSFERASE TRANSFORMYLASE, PURINE BIOSYNTHESIS, ATP-GRASP	OXIDOREDUCTASE LYSINE BIOSYNTHESIS, ALPHA- AMINOADIPATE PATHWAY, 2 SACCHAROPINE REDUCTASE, DEHYDROGENASE	OXIDOREDUCTASE OXIDOREDUCTASE, OXIDOREDUCTASE, NAD	TRANSFORMING GROWTH FACTOR
Coumpound		NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130), CHAIN: A;	ALPHA-I SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES I-130); CHAIN: B;	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN: A;		N-(I-D-CARBOXYLETHYL)- L-NORVALINE DEHYDROGENASE; CHAIN: NULL;	GLYCERALDEHYDE-3- PHOSPHATE DEHYDROGENASE; CHAIN: P, R, O, Q;	PHOSPHORIBOSYLGLYCINA MIDE FORMYLTRANSFERASE 2; CHAIN: A, B;	SACCHAROPINE REDUCTASE; CHAIN: A;	L-ALANINE DEHYDROGENASE; CHAIN: A;	BONE MORPHOGENETIC
SeqFold score												
PMF score		0.57	0.86	<u>-</u>	0.45		0.17	60.0	0.05	-	0.49	_
Verify score		-0.12	0.49	0.28	-0.09		-0.18	-0.1	60'0-	1.04	-0.24	0.14
PSI- BLAST		9.00E-15	3.20E-17	1.40E-18	6.40E-17		1.60E-05	0.0069	0.0054	4.80E-58	1.30E-61	9.60E-48
End		106	103	107	66		594	570	599	924	460	213
Start AA		21	20	20	23		481	480	482	481	25	112
Chain ID		V	<	₹	A			۵۰	4	<	V	
PDB ID		1qau	Iqav	1qlc	3pdz		15g6	1cD	leyz	6 <u>H</u>	1pjc	16m
SEQ DO:		838	838	838	838		840	840	840	840	840	842

					 -				
PDB annotation	BMP-7; MORPHOGEN, TRANSFORMING GROWTH FACTOR, CYTOKINE, BONE, 2 CARTILAGE, GLYCOPROTEIN	TRANSFORMING GROWTH FACTOR OSTEOGENIC PROTEIN-1, HOP-1, BMP-7, MORPHOGEN, TRANSFORMING GROWTH FACTOR, CYTOKINE, BONE, 2 CARTILAGE, GLYCOPROTEIN				CYTOKINE CYTOKINE, BONE MORPHOGENETIC PROTEIN, CYSTIN- KNOT, TGFB- 2 FAMILY	HYDROLASE TETRATRICOPEPTIDE, TRP, HYDROLASE, PHOSPHATASE, PROTEIN-PROTEIN INTERACTIONS, TPR, 2 SUPER-HELIX, X-RAY STRUCTURE	HYDROLASE TETRATRICOPEPTIDE, TRP, HYDROLASE, PHOSPHATASE, PROTEIN-PROTEIN INTERACTIONS, TPR, 2 SUPER-HELIX, X-RAY STRUCTURE	HYDROLASE TETRATRICOPEPTIDE, TRP, HYDROLASE, PHOSPHATASE, PROTEIN-PROTEIN INTERACTIONS, TPR, 2 SUPER-HELIX, X-RAY STRUCTURE
Coumpound		BONE MORPHOGENETIC PROTEIN-7; CHAIN: NULL;	GROWTH FACTOR TRANSFORMING GROWTH FACTOR-BETA TWO (TGF- B2) 2TGI 3	GROWTH FACTOR TRANSFORMING GROWTH FACTOR-BETA TWO (TGF- B2) 2TGI 3	GROWTH FACTOR TRANSFORMING GROWTH FACTOR-BETA TWO (TGF- B2) 2TGI 3	BONE MORPHOGENETIC PROTEIN 2 (BMP-2); CHAIN: A;	SERINE/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;	SERINE/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;	SERINE/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;
SeqFold score		104.28	67.81				·		
PMF				0.81	9.0	-	0.25	0.11	0.8
Verify score				2 .0	0.11	0.05	0.16	0.23	0.47
PSI- BLAST		9.60E-48	1.80E-38	1.60E-36	1.80E-38	4.80E-48	0.00013	1.60E-07	1.40E-05
End		214	214	213	213	213	605	803	1103
Start		112	100	102	112	Ξ	430	999	992
Chain ID						A			
PDB ID		1bm p	2tgi	2tgi	2tgi	3bm p	lal7	1817	1817
SEQ ID	Ö	842	842	842	842	842	843	843	843

9	4			1	Dog	Varify	DME	Seaffold	Commonnd	PDB annotation
) (1)	<u> </u>	E A	AA	AA AA	BLAST	score	score	score		
2									CHAIN: A, B; PTS1- CONTAINING PEPTIDE; CHAIN: C, D;	PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT
843	1fch	<	445	804	3.60E-09	0.29	0.62		PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTSI- CONTAINING PEPTIDE; CHAIN: C, D;	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT
843	1fch	<	452	669	1.30E-20	0.06	-0.15		PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTSI- CONTAINING PEPTIDE; CHAIN: C, D;	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT
843	1fch	4	576	842	1.10E-12	0.02	66'0		PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTSI- CONTAINING PEPTIDE; CHAIN: C, D;	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT
843	1fyn	<	193	242	1.80E-07	0.23	4.0		PHOSPHOTRANSFERASE FYN: CHAIN: A; 3BP-2; CHAIN: B;	TRANSFERASE PROTO-ONCOGENE TYROSINE KINASE; PROTO- ONCOGENE, TRANSFERASE, PHOSPHORYLEN KINASE, 2 PHOSPHORYLATION, SH3 DOMAIN. 3 COMPLEX (PHOSPHOTRANSFERASE/PEPTIDE)
843	1gbr	<	181	249	1.40E-06	0.74	0.59		SIGNAL TRANSDUCTION PROTEIN GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2, N-TERMINAL IGBR 3 SH3 DOMAIN) COMPLEXED WITH SOS-A PEPTIDE IGBR 4 (NMR, 29 STRUCTURES) IGBR 5	
843	11 Sk	<	193	288	1.80E-05	0.09	0.28		P56=LCK= TYROSINE KINASE; ILCK 7 CHAIN: A; ILCK 8 TAIL PHOSPHOPEPTIDE TEGQ(PHOSPHO)YQPQPA; ILCK 14 CHAIN: B; ILCK 15	COMPLEX (KINASE/PEPTIDE)
843	Inlo	C	193	242	1.80E-07	4.0	0.29		C-SRC, CHAIN: C, NL1 (MN7-	COMPLEX (TRANSFERASE/PEPTIDE)

					ż ,		7			T @ ^
r DD alliforation	SRC, SH3 DOMAIN, LIGANDS, NON- PEPTIDE ELEMENTS, 2 COMPLEX (TRANSFERASE/PEPTIDE)		TYROSINE KINASE I YKUSINE KINASE-INHIBITOR COMPLEX, DOWN-REGULATED KINASE, 2 ORDERED ACTIVATION LOOP	PROTEIN TRANSPORT HELIX-TORN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT	PROTEIN TRANSPORT HELLX-LUKN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT		TRANSFERASE HCK; SH3, PROTEIN TYROSINE KINASE, SIGNAL TRANSDUCTION, 2 TRANSFERASE	IMMUNOGLOBULIN IMMUNOGLOBULIN	IMMUNOGLOBULIN, FAB IMMUNOGLOBULIN, FAB FRAGMENT, HUMANISATION	COMPLEX (VIRAL. CAPSID/IMMUNOGLOBULIN) HIV-1 CA, HIV CA, HIV P24, P24; FAB, FAB LIGHT CHAIN, FAB HEAVY CHAIN COMPLEX (VIRAL CAPSID/IMMI NOGI, OBULIN), HIV.
Coumpound	MN2-MN1-PLPPLP); CHAIN: N;	PHOSPHOTRANSFERASE PHOSPHATIDYLNOSITOL 3- KINASE (P85-ALPHA SUBUNIT, IPNI 3 SH3 DOMAIN) (NMR, MINIMIZED AVERAGE STRUCTURE) IPNI 4	HAEMATOPOETIC CELL KINASE (HCK); CHAIN: A;	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	PHOSPHOTRANSFERASE FYN PROTO-ONCOGENE TYROSINE KINASE (E.C.2.7.1.112) 1SHF 3 (SH3 DOMAIN) 1SHF 4	HEMATOPOIETIC CELL KINASE; CHAIN: NULL;	2E8 (IGG1=KAPPA=) ANTIBODY; CHAIN: L, H, M, P;	ANTIBODY CTM01; CHAIN: L, H;	HUMAN IMMUNODEFICIENCY VIRUS TYPE I CAPSID CHAIN: A. B; ANTIBODY FAB25.3 FRAGMENT; CHAIN: H. K. L.
SeqFold										
PMF score		0.17	0.17	0.15	60.0	0.54	0.89	-0.06	0.03	0.24
Verify		0.54	0.09	90.0	-0.03	0.38	0.45	0.08	-0.18	0.05
PSI- BLAST		9.00E-10	3.60E-07	1.80E-07	8.00E-06	9.00E-08	9.00E-06	6.40E-51	4.80E-53	4.80E-63
End		249	282	573	1101	242	255	414	599	220
Start		182	193	418	949	188	193	236	419	21
Chain ID			«	A	A	<		H	ж	H
PDB		1pnj	lqcf	Iqqe	Idde	lshf	4hck	1.20 E+09	lac6	1afv
SEQ ID	NO:	843	843	843	843	843	843	846	846	846

		_												_				AL	 !				_				LEA,										<u>.</u>		O V C	7. AB
	CAPSID PROTEIN, 2 P24	INSECT IMMUNITY INSECT	IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	NICECT IMMINITY INSECT	MAN MINISTER TENTING	HOMOPHI I, C. 3-BINDING,	PISTON INDIAN INTENT	INSECT INTRODUCTOR	IMMONII I, EFS-BIIADIIAG,	HOMOT HILLS ADMINISTER	NSECT IMMONTED INSECT	IMMONII Y, LPS-BINDING, HOMOPHII IC ADHESION	NSECT INAMINITY INSECT	PARTITY 1 PS-BINDING.	HOMOPHILIC ADHESION	RECEPTOR RECEPTOR, SIGNAL	TRANSPILCER OF 11-6 TYPE	CVTOVINES THIRD 2 N. TERMINAL	DOMAIN TRANSMEMBRANE.	GLYCOPROTEIN	IMMUNE SYSTEM	IMMUNOGLOBULIN, IMMUNE	SYSTEM			IMMUNOGLOBULIN	IMMUNOGLOBULIN, FAB COMPLEY,	IDIOTOPE, ANTI-IDIOLOPE					IMMUNE SYSTEM	IMMUNOGLOBULIN, 1001,	IMMUNOGLOBULIN, IGGI;	IMMUNOGLOBULIN, 1GG1 FAB	FRAGMENT, CROSS-REACTIVITY	HIVI PROTEASE, ENZYME 2	INHIBITION, IMMORACE CONCENT	IMMINE SYSTEM ANTI-FROM
Coumpound	0						1			1					-		_				MONOCI ONAL ANTIBODY		NAL	ANTIBODY MRK-16 (HEAVY	-			HEAVY CHAIN V REGIONS;	CHAIN: B; IG HEAVY CHAIN	V REGIONS; CHAIN: C; IG	HEAVY CHAIN V REGIONS;		9691 AQC	(LIGHT CHAIN); CHAIN: L;	3DY 1696	(VARIABLE HEAVY CHAIN);	CHAIN: H; IGG1 ANTIBODY	1696 (CONSTANT HEAVY	VIN: 1;	DV LIGHT
Conmi		TIENOT INI. CUAIN: A B.	HEMOLIN, CH.		HEMOLIN; CHAIN: A, B;			HEMOLIN; CHAIN: A, B;			HEMOLIN; CHAIN: A, B;		10 14 107 1011	HEMOLIN; CHAIN: A, B,		Chizo. CHANI MIII	GF130; CFIAUN				MONOCION	MRK-16 (LIGHT CHAIN);	CHAIN: A. C.	ANTIBODY M	CHAIN); CHAIN: B, D;	IG HEAVY CHAIN V	REGIONS; CHAIN: A; 1G	HEAVY CHA	CHAIN: B; IG	V REGIONS;	HEAVY CHA	CHAIN: D.	IGGI ANTIBODY 1696	(LIGHT CHA)	IGG1 ANTIBODY 1696	(VARIABLE I	CHAIN: H; IG	1696 (CONST	CHAIN); CHAIN: I;	EAR ANTIRODY LIGHT
SeqFold		0000	76.661																																					
PMF					_			0.54			99.0			0.21			-0.05				200	5.0				-0.13							-0.19							
Verify score					0.28			0.29			0.15			0.14			0.07				76.0	0.30				0.17							0.03	_						70.0
PSI- BLAST			5.40E-62		5.40E-62			1.60E-31			7.20E-48			3.20E-22			1.40E-12				2 200 (4	3.20E-04				1 60F-49							6.40E-23			-				2 20E 40
End AA			503		503			493			404			713			597			_	3	077				412	! : 					_	603				-			717
Start			134		135	<u>.</u>		138))		77	ì		323			208					7.1				235	2						523							23.0
Chain ID			A		▼	:		4	:		4	ć		<								æ				a	۵						_	í						
PDB m			1bih		1hih	5		1. High	-		4.4			1bih			16,8	,				15m				100	ڍ						1017			_				
SEQ ID	Ö		846		846			846	2		24.0	0.50		846			846					846				946	0 0						846	2						1

je Š	5			End	PSI-	Verity	PML	SedFold	Conmbonna	
		E	¥¥	¥	BLAST	score	score	score		
_									CHAIN; CHAIN: L; FAB ANTIBODY HEAVY CHAIN; CHAIN: H:	3F4; ANTI-PRION FAB 3F4 ANTI-PRION ANTIBODY, FAB 3F4
846 1	lcs6	¥	135	503	3.60E-59	80:0	-		AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
846	1cs6	A	138	503	4.80E-45	0.24			AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
846	lcs6	<	2	321	1.60E-47	-0.14	0.43		AXONIN-1; CHAIN: A:	CELL ADHESION NEURAL CELL ADHESION
846	1cs6	<	20	412	1.10E-37	0.02	-0.14		AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
846	1cs6	∢	226	592	1.80E-44	0.23	96.0		AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
846	1cs6	<	236	602	6.40E-42	0.13	-		AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
846	1cs6	V	24	404	1.60E-53	0.07	0.58		AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
846	1cs6	4	320	713	3.20E-33	0.07	0.51		AXONIN-1; CHAIN: A;	CELL ADHESION NEURAL CELL ADHESION
╁	10.00	ر	125	320	1 80F-30	0 15	0.94		FIBROBLAST GROWTH	GROWTH FACTOR/GROWTH FACTOR
	<u>.</u>	ر ر	<u></u>	3		<u>}</u>			FACTOR 2; CHAIN: A, B;	RECEPTOR FGF, FGFR,
								_	FIBROBLAST GROWTH	TEANSDICTION 2 DIMERIZATION.
_									FACTOR RECEPTOR 1; CHAIN: O. D:	GROWTH FACTOR/GROWTH FACTOR
				_						RECEPTOR
846	levs	C	430	109	3.20E-21	0.07	0.22		FIBROBLAST GROWTH	GROWTH FACTOR/GROWTH FACTOR
	3)	<u>}</u>	<u>:</u>					FACTOR 2; CHAIN: A, B;	RECEPTOR FGF, FGFR,
_									FIBROBLAST GROWTH	IMMUNOGLOBULIN-LIKE, SIGNAL
_									FACTOR RECEPTOR 1;	TRANSDUCTION, 2 DIMENZATION, CROWNTH FACTOR
									CHAIN: C, D;	RECEPTOR
846	levs	C	135	320	1.80E-32	0.22	0.99		FIBROBLAST GROWTH	GROWTH FACTOR/GROWTH FACTOR
	?	1	-))					FACTOR 2; CHAIN: A, B;	RECEPTOR FGF, FGFR,
									FIBROBLAST GROWTH	IMMUNOGLOBULIN-LIKE, SIGNAL
_									PACTOR RECEPTOR 1;	GROWTH FACTOR/GROWTH FACTOR
_		· · · -								RECEPTOR
846	158	6	430	109	8.00E-21	0.26	-0.03		FIBROBLAST GROWTH	GROWTH FACTOR/GROWTH FACTOR
	2	ì							FACTOR 2; CHAIN: A, B;	RECEPTOR FGF, FGFR,
									FIBROBLASI GROWIH	TRANSPICTION 2 DIMERIZATION.

PDB annotation	GROWTH FACTOR/GROWTH FACTOR RECEPTOR	R; VIRUS/VIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, IAIN: ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	R; VRUS/VIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, IAIN: ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	1; VIRUS/VIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, IAIN: ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	IMMUNE SYSTEM FC IGG PHAGE IN; DISPLAY PEPTIDE RED	HAIN: IMMUNE SYSTEM IMMUNOGLOBULIN FOLD, ANTIBODY, IGM, FV	EGG- BETA-N-ACETYLMURAMIDASE C; SINGLE-DOMAIN ANTIBODY, TURKEY EGG-WHITE LYSOZYME, 2 ANTIBODY-PROTEIN COMPLEX, SINGLE-CHAIN FV FRAGMENT	 		I GROWTH FACTOR/GROWTH FACTOR S, C, D; RECEPTOR FGF2; FGFR2; I IMMUNOGLOBULIN (IG)LIKE
Coumpound	CHAIN: C, D;	POLIOVIRUS RECEPTOR; CHAIN: R; VP1; CHAIN: 1; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	POLIOVIRUS RECEPTOR; CHAIN: R; VPI; CHAIN: 1; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	POLIOVIRUS RECEPTOR; CHAIN: R; VPI; CHAIN: 1; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	IMMUNOGLOBULIN LAMBDA HEAVY CHAIN; CHAIN: A, B; ENGINEERED PEPTIDE; CHAIN: E, F;	IGM MEZ IMMUNOGLOBULIN; CHAIN: L; IGM MEZ IMMUNOGLOBULIN; CHAIN: H;	SCEV FRAGMENT 1F9; CHAIN: A, B; TURKEY EGG- WHITE LYSOZYME C; CHAIN: X, Y;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, FIBROBLAST GROWTH
SeqFold score										
PMF score		0.65	0.21	0.01	0.05	60.0-	-0.03	6.0	0.78	96.0
Verify score		-0.3	-0.31	-0.4	-0.27	60.0	0.14	0.34	0.37	0.07
PSI- BLAST		7.20E-36	1.10E-38	3.60E-44	8.00E-40	3.20E-38	6.40E-42	1.80E-31	1.30E-24	1.40E-34
End		404	503	319	411	130	206	319	\$03	320
Start		141	228	28	234	21	21	142	324	135
Chain ID		~	~	<u>«</u>	¥	н	∢	V	¥	ப
PDB ID		Idgi	1dgi	Idgi	1dn2	1 dq1	ldzb	lepf	lepf	lcv2
SEQ ID NO:		846	846	846	846	846	846	846	846	846

PDB annotation	DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTORIGROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1; FGFR1; IMMUNOGLOBULN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1, FGFR1, IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	IMMUNE SYSTEM FC-EPSILON RI- ALPHA; IMMUNOGLOBULIN FOLD,
Coumpound	FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 1; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR I; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C. D;	HIGH AFFINITY IMMUNOGLOBULIN
SeqFold score								
PMF score		0.17	0.94	0.94	0.07	69:0	0.07	0.29
Verify score		0.15	0.23	0.27	0.15	0.11	-0.07	60.0
PSI- BLAST		7.20E-23	5.40E-33	3.60E-24	1.60E-19	1.30E-30	6.40E-20.	7.20E-28
End		224	322	505	109	320	601	322
Start		33	135	329	430	135	430	136
Chain ID		ല	O	ט	Ü	ပ	U	<
PDB ID		lev2	lev2	lev2	lev2	levt	levi	1f2q
SEQ	Ž	846	846	846	846	846	846	846

SEQ ID	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
									EPSILON RECEPTOR CHAIN: A;	GLYCOPROTEIN, RECEPTOR, IGE- BINDING 2 PROTEIN
846	1f4w	н	21	220	1.60E-62	0.02	0		ANTIBODY S-20-4, FAB FRAGMENT, LIGHT CHAIN; CHAIN: L; ANTIBODY S-20-4, FAB FRAGMENT, HEAVY CHAIN; CHAIN: H	IMMÜNE SYSTEM ANTI- CARBOHYDRATE ANTIBODY
846	l f6a	«	132	322	1.10E-29	0.17	0.18		HIGH AFFINITY IMMUNOGLOBULIN EPSILON RECEPTOR CHAIN: A; IG EPSILON CHAIN C REGION; CHAIN: B, D;	IMMUNE SYSTEM HIGH AFFINITY IGE-FC RECEPTOR, FC(EPSILON) IGE- FC, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN, RECEPTOR, IGE- BINDING 2 PROTEIN, IGE ANTIBODY, IGE-FC
846	1f6f	В	524	715	9.00E-15	0.05	0.03		PLACENTAL LACTOGEN; CHAIN: A; PROLACTIN RECEPTOR; CHAIN: B, C;	HORMONE/GROWTH FACTOR/HORMONE RECEPTOR 4- HELICAL BUNDLE, ALPHA HELICAL BUNDLE, TERNARY COMPLEX, FN 2 III DOMAINS, BETA SHEET DOMAINS, CYTOKINE-RECEPTOR COMPLEX
846	1fc2	Ω	234	411	1.10E-39	-0.03	0.07		IMMUNOGLOBULIN IMMUNOGLOBULIN FC AND FRAGMENT B OF PROTEIN A COMPLEX 1FC2 4	
846	lfcg	4	143	319	1.10E-24	-0.16	0.23		FC RECEPTOR FC(GAMMA)R11A; CHAIN: A;	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32
846	1 finh	V.	512	208	3.60E-17	0.17	0.52		FIBRONECTIN; CHAIN: A;	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING
846	<u>।</u> मा	⋖	139	319	1.80E-22	-0.02	0.36		LOW AFFINITY IMMUNOGLOBULIN GAMMA FC REGION CHAIN: A;	IMMUNE SYSTEM RECEPTOR BETA SANDWICH, IMMUNOGLOBULIN- LIKE, RECEPTOR
846	lhnf		141	305	1.60E-21	-0.31	0.04		T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) IHNF 3	
846	lhng	∢	143	319	3.60E-27	0.03	0.05		T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) IHNG 3	
846	ligt	В	21	412	0	-0.16	0.11		IGG2A INTACT ANTIBODY - MAB231; CHAIN: A, B, C, D	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN V REGION C

SEQ	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
ÿ										REGION, IMMUNOGLOBULIN
846	ligy	æ	13	412	3.20E-95			110.16	IGGI INTACT ANTIBODY MAB61.1.3; CHAIN: A, B, C, D	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION
846	ligy	В	21	411	3.20E-95	-0.05	0.24		IGGI INTACT ANTIBODY MAB61.1.3; CHAIN: A. B, C, D	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION
846	litb	В	225	503	5.40E-29	0.21	0.49		INTERLEUKIN-I BETA; CHAIN: A; TYPE I INTERLEUKIN-I RECEPTOR; CHAIN: B;	COMPLEX— (IMMUNOGLOBULIN/RECEPTOR) IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, RECEPTOR, 2 SIGNAL, COMPLEX (IMMUNOGLOBULIN/RECEPTOR)
846	1kb5	=	235	413	1.10E-49	0.11	-0.17		KB5-C20 T-CELL ANTIGEN RECEPTOR; CHAIN: A, B; ANTIBODY DESIRE-1; CHAIN: L, H;	COMPLEX (IMMUNOGLOBULIN/RECEPTOR) TCR VAPLHA VBETA DOMAIN; T-CELL RECEPTOR, STRAND SWITCH, FAB, ANTICLONOTYPIC, 2 (IMMUNOGLOBULIN/RECEPTOR)
846	Ilmk	<	21	206	3.20E-39	-0.23	0.01		IMMUNOGLOBULIN ANTI- PHOSPHATIDYLINOSITOL SPECIFIC PHOSPHOLIPASE C DIABODY ILMK 3 SYNONYMS: L5MK16 DIABODY, SINGLE-CHAIN FV DIMER ILMK 4	
846	Ппа	工	21	220	4.80E-62	-0.05	0.22		IMMUNOGLOBULIN ANTIGEN-BINDING FRAGMENT (FAB) (IGG2B, KAPPA) IMAM 3	
846	1mc o	II	50	411	1.60E-93	-0.17	0.04		IMMUNOGLOBULIN IMMUNOGLOBULIN GI (IGGI) (MCG) WITH A HINGE DELETION IMCO 3	
846	lmc o	ж	24	416	1.60E-93			123.28	IMMUNOGLOBULIN IMMUNOGLOBULIN GI (IGG1) (MCG) WITH A HINGE DELETION IMCO 3	
846	1mc P	I	21	210	8.00E-46	-0.26	0.06		IMMUNOGLOBULIN IMMUNOGLOBULIN FAB FRAGMENT (MC/PC\$603)	

													Τ						T	7
PDB annotation	CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, FYTRACELLULAR MATRIX, 2	HEPARIN-BINDING, GLYCOPROTEIN			COMPLEX (IMMUNORECEPTOR/IMMUNOGLOBU	(IMMUNORECEPTOR/IMMUNOGLOBU	IMMUNOGLOBULIN IMMINOGLOBULIN.	IMMUNOGLOBULIN VARIABLE	HEAVY (VH) DOMAIN, VARIABLE IIGHT (VL) ANTIBODY FRAGMENT,	MULTIVALENT ANTIBODY,	DIABOD I, DOMALIN 2 3 WALL INC.	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2	CTRICTINAL PROTEIN INTEGRIN.	HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2	PROTEIN	IMMUNE STSTEM FAB, PORA, NEISSERIA MENINGITIDIS, PORIN		IMMUNOGLOBULIN IMMUNOGLOBULIN, SINGLE-CHAIN	FV, AN II-CARCINOEMBA FOR CE	STRUCTURAL PROTEIN TENASCIN,
Coumpound	IMCP 4 FIBRONECTIN, CHAIN: NULL;		IMMUNOGLOBULIN IGG JEL 103 FAB FRAGMENT COMPI FXED WITH IMRD 3	INOSINE-5'-DIPHOSPHATE IMRD 4	NIS ALPHA-BETA T-CELL RECEPTOR; CHAIN: A, B, C,	D; H57 FAB; CHAIN: E, F, G, H	NIG9 (IGGI=LAMBDA=);	SINGLE-CHAIN ANTIBODY	FRAGMENT; CHAIN: A, C;			INTEGRIN BETA-4 SUBUNIT; CHAIN: A. B;	TIMIET A TOTAL A CITE INTE	CHAIN: A, B;		ANTIBODY; CHAIN: H, L; PROTEIN G-PRIME; CHAIN: A: MAJOR OUTER	MEMBRANE PROTEIN P1.16; CHAIN: P:	MFE-23 RECOMBINANT ANTIBODY FRAGMENT;	CHAIN: A;	TENASCIN; CHAIN: A, B;
SeqFold score																	<u></u>			
PMF	0.15		90:0-		-0.05		0.13	0.04	5			0.21		0.24		-0.01		0.25		0.19
Verify	-0.03		0.05		0.05		0.14	11	<u>:</u>			0.08		0.0		0.12		0.24		0.04
PSI- BLAST	3.60E-21		9.60E-64		4.80E-62		1.60E-63	2 20E 43	3.20E-43			5.40E-15		1.80E-23		3.20E-63		8.00E-40		5.40E-19
End	715		220		220		220	200	007			599		715		220		211		713
Start	512		21		21		21	ć	77			449		508		21		21		512
Chain ID			H		i.		H		∢ .			V		<		=		<		V
PDB CI	1mfn		Imrd		lnfd		Ingp		dbu [1983		1483		1qkz		1qok		lar4
SEQ	%O:		846		846		846		846			846		846		846		846		846

SEQ ID	PDB ID	Chain 1D	Start AA	End	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	LDB alliciation
Ö										FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN
846	lvca	4	137	319	3.60E-24	0.19	0.74		HUMAN VASCULAR CELL ADHESION MOLECULE-1; 1VCA 4 CHAIN: A, B; 1VCA 5	CELL ADHESION PROTEIN VOAVA- DI.2: IVCA 6 IMMUNOGLOBULIN SUPERFAMILY, INTEGRIN-BINDING IVCA 15
846	lwej	π	236	413	8.00E-50	0.09	-0.14		E8 ANTIBODY; CHAIN: L, H; CYTOCHROME C; CHAIN: F;	COMPLEX (ANTIBOD FELL CINCIA TRANSPORT) FAB E8, CYT C, ANTIGEN: IMMUNOGLOBULIN, IGGI KAPPA, FAB FRAGMENT, HORSE 2 CYTOCHROME C, COMPLEX (ANTIBODY/FELECTRON TRANSPORT)
846	1 yuh	H	419	602	8.00E-53	0	-0.15		FAB FRAGMENT; CHAIN: NULL;	IMMUNOGLOBULIN AN 11- NITROPHENOL, LAMBDA LIGHT CHAIN, IMMUNOGLOBULIN
846	1zxq		135	326	1.80E-34	0	60.0		INTERCELLULAR ADHESION MOLECULE-2; CHAIN: NULL;	CELL ADHESION ICAM'2, IMMUNOGLOBULIN FOLD, CELL ADHESION, GLYCOPROTEIN, 2 TRANSMEMRANE, REPEAT, SIGNAL
846	2dli	∢	134	320	7.20E-28	0.29	0.09		MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	IMMUNE SYSTEM PS NATURAL KILLER CELL RECEPTOR; KIR, NATURAL KILLER RECEPTOR, INHIBITORY RECEPTOR, 2 IMMUNOGLOBULIN
846	2fbj	H	21	210	3.20E-50	0.12	0.19		IMMUNOGLOBULIN 1G*A FAB FRAGMENT (1539) (GALACTAN-BINDING) 2FBJ 3	dOTTO TO TO TO THE TAXABLE TO THE TA
846	2fcb	<	135	323	7.20E-36	0.13	0.81		FC GAMMA RIIB; CHAIN: A;	IMMUNE SYSTEM CD32; RECEFTOR, FC, CD32, IMMUNE SYSTEM TAXABLE SYSTEM CD32: RECEPTOR
846	2fcb	< -	226	410	9.00E-29	-0.04	0.21		FC GAMMA RIIB; CHAIN: A; IMMUNOGLOBULIN FAB	IMMUNE SYSTEM FC, CD32, IMMUNE SYSTEM
846	8 tab	<	234	704	9.00E-20	CC:			FRAGMENT FROM HUMAN IMMUNOGLOBULIN IGGI (LAMBDA, HIL) 8FAB 3	
847	1dp.1	3	299	480	1.60E-09	0.05	0.11		SYNTAXIN BINDING PROTEIN 1, CHAIN: A;	ENDOCYTOSIS/EXOCYTOSIS NSECI, PROTEIN-PROTEIN COMPLEX,

SeqFold Coumpound PDB annotation score	SYNTAXIN 1A; CHAIN: B; MULTI-SUBUNIT	CHITINASE I: CHAIN: A: HYDROLASE BETA-ALPHA BARREL		CHITINASE B: CHAIN: A, B; HYDROLASE HYDROLASE, CHITIN DEGRADATION	CHITINASE B; CHAIN: A, B; HYDROLASE HYDROLASE, CHITIN DEGRADATION	CHITINASE A; CHAIN: A; HYDROLASE BETA-ALPHA (TIM) BARREL	CHITINASE A; CHAIN: A; HYDROLASE BETA-ALPHA (TIM) BARREL	HYDROLASE (GLUCOSIDASE) ENDO- BETA-N-	ACETYLGLUCOSAMINIDAS E H, ENDO H (E.C.3.2.1.96)	HYDROI AGE/GI HONGIDA GE) ENDO-BETA-N-	ACETYLGLUCOSAMINIDAS	E FI (E.C.3.2.1.96) ZEBN 3 ·	ENDO F1) 2EBN 4	HYDROLASE(GLUCOSIDASE	ACETYLGLUCOSAMINIDAS	E F1 (E.C.3.2.1.96) 2EBN 3	(ENDOGE YCOSIDASE FI, ENDO FI) 26BN 4	77.33 DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN	(HOMEODOMAIN) MUTANT	WITH CYS 39 1AHD 3	REPLACED BY SER (C39S)	COMPLEA WILL DIVA (NIMR,	1AHD 4 16 STRUCTIRES)
PMF S		-	-0.09	_			-	0.13		90.0					0.77				7					
Verify		0.35	0.19	0.47	0.51	0.41	0.33	-0.15		350	3.0		_		-0.3									
PSI- BLAST		4 ROF-63	1.60E-10	1.10E-81	1.60E-59	3.60E-79	9.60E-70	4.80E-05		7 20E 51	10-707:/	_			0.0067				7.20E-33				_	
End AA		707	367	265	266	281	260	146		200					Ξ				351					
Start		2	323	-	2	1	4	4		-	•				4				284					_
Chain ID		A	: ×	4	4	4	<												d.					
PDB ID		1426	1dqc	1.00 E+15	1.00 E+15	ledq	ledq	ledt		2.hn					2ebn				lahd					_
SEQ ID		848	848	848	848	848	848	848		87.8	2				848				849				_	_

CEO	PDR	Chain	Start	End	-ISd	Verify	PMF	SeqFold	Coumpound	PDB annotation
i e ș	<u>e</u>	<u>a</u>	AA	VV	BLAST	score	score	score		
2									ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 1 AHD 3	
									REPLACED BY SER (C39S)	
									IAHD 4 16 STRUCTURES)	
849	lahd	Ы	286	351	7.20E-33	-0.38	1-1		DNA-BINDING PROTEIN	
 }									ANTENNAPEDIA PROTEIN	
									(HUMEUDOMAIN) MOTANT WITH CYS 39 IAHD 3	
									REPLACED BY SER (C39S)	
									COMPLEX WITH DNA (NMR,	
									1AHD 4 16 STRUCTURES)	
849	1672	<	274	346	5.40E-32			62.11	HOMEOBOX PROTEIN HOX-	PROTEIN/DNA HOMEODOMAIN, DNA,
									B1; CHAIN: A; PBX1; CHAIN:	COMPLEX, DNA-BINDING FROI EIN. PROTEIN/DNA
									CHAIN: E;	
849	1672	V	286	346	5.40E-32	-0.08	_		HOMEOBOX PROTEIN HOX-	PROTEIN/DNA HOMEODOMAIN, DNA,
									BI; CHAIN: A; PBXI; CHAIN:	COMPLEX, DNA-BINDING PROTEIN,
									B; DNA CHAIN: D; DNA CHAIN: E;	PROTEIN/DINA
840	1,48;	Ψ	285	342	\$ 40E-31			65.54	ULTRABITHORAX	TRANSCRIPTION/DNA
Ì	5	:	-	<u>.</u>					HOMEOTIC PROTEIN IV;	ULTRABITHORAX; PBX PROTEIN,
									CHAIN: A; HOMEOBOX	DNA BINDING, HOMEODOMAIN,
					_	_			PROTEIN EXTRADENTICLE;	HOMEOTIC PROTEINS,
									C; DNA (5'- CHAIN: D;	
648	158i	4	286	342	5.40E-31	-0.11	0.99		ULTRABITHORAX	TRANSCRIPTION/DNA
									HOMEOTIC PROTEIN IV;	ULTRABITHORAX; PBX PROJEIN;
							-		CHAIN: A; HOMEOBOA	HOMEOTIC PROTEINS.
									CHAIN: B. DNA (5'- CHAIN:	DEVELOPMENT, 2 SPECIFICITY
									C; DNA (5'- CHAÎN: D;	
849	140s	٧	35	238	7.20E-11	0.15	-0.19		NICOTINATE	TRANSFERASE DINUCLEOTIDE-
		_							MONONUCLEOIIDE:3,8- CHAIN: A;	BINDING MOTIF, FINOSI HONBOSTE TRANSFERASE
849	lftz		283	351	1.30E-27			68.94	DNA-BINDING FUSHI	
								_	TARAZU PROTEIN (HOMEODOMAIN) (NMR, 20	
				4						

EQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
<u>e ë</u>	<u>e</u>	e	¥	¥¥	BLAST	score	score	score		
									STRUCTURES) IFTZ 3	
849	l ftz		284	337	1.30E-27	0.31	0.84		DNA-BINDING FUSHI TARAZU PROTEIN (HOMEODOMAIN) (NMR, 20 STRUCTURES) IFTZ 3	
849	 Oct	ပ	205	344	7.20E-34	-0.2	0.12		DNA-BINDING PROTEIN OCT-1 (POU DOMAIN) IOCT 3	
849	Isan		290	351	5.40E-30			72.13	DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 ISAN 3 REPLACED BY SER AND RESIDUES 1-6 DELETED (C398,DEL 1-6) ISAN 4 (NMR,	
849	Isan		291	337	1.40E-26	-0.21			DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 ISAN 3 REPLACED BY SER AND RESIDUES 1-6 DELETED (C39S,DEL 1-6) ISAN 4 (NMR, 20 STRUCTURES) ISAN 5	
849	lsan		292	351	5.40E-30	-0.2	-		DNA-BINDING PROTEIN ANTENNAPEDIA PROTEIN (HOMEODOMAIN) MUTANT WITH CYS 39 ISAN 3 REPLACED BY SER AND RESIDUES 1-6 DELETED (C39S,DEL 1-6) ISAN 4 (NMR, 20 STRUCTURES) ISAN 5	
849	9ant	٧	289	337	1.30E-27	0.11	-		ANTENNAPEDIA PROTEIN; CHAIN: A, B; DNA; CHAIN: C, D, E, F;	COMPLEX (DNA-BINDING PROTEIN/DNA) HD; HOMEODOMAIN, COMPLEX (DNA-BINDING PROTEIN/DNA)
849	9ant	4	289	344	9.00E-30			90.69	ANTENNAPEDIA PROTEIN; CHAIN: A, B; DNA; CHAIN: C, D, E, F;	COMPLEX (DNA-BINDING PROTEIN/DNA) HD; HOMEODOMAIN, COMPLEX (DNA-BINDING PROTEIN/DNA)
849	9ant	4	289	344	9.00E-30	0.27			ANTENNAPEDIA PROTEIN;	COMPLEX (DNA-BINDING

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
Ö									CHAIN: A, B; DNA; CHAIN: C, D, E, F;	PROTEIN/DNA) HD; HOMEODOMAIN, COMPLEX (DNA-BINDING PROTEIN/DNA)
851	la6a	В	33	157	3.20E-66			54.43	HLA-DR3; CHAIN: A, B; CLIP; CHAIN: C;	COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN) MHC GLYCOPROTEIN, COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN)
851	1a6a	B	34	147	3.20E-66	-0.32	0.93		HLA-DR3; CHAIN: A, B; CLIP; CHAIN: C;	COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN) MHC GLYCOPROTEIN, COMPLEX (TRANSMEMBRANE/GLYCOPROTEIN)
851	laqd	В	33	147	6.40E-69	-0.33	0.63		HLA-DRI CLASS II HISTOCOMPATIBILITY PROTEIN; CHAIN: A, B, D, E, G, H, J, K; HLA-A2; CHAIN: C, F, I, L;	COMPLEX (MHC PROTEINAN LIGEN) DRA, DRB1 01010; COMPLEX (MHC PROTEIN/ANTIGEN), HISTOCOMPATIBILITY ANTIGEN
851	laqd	В	33	157	6.40E-69			55.9	HLA-DR1 CLASS II HISTOCOMPATIBILITY PROTEIN; CHAIN: A, B, D, E, G, H, J, K; HLA-A2; CHAIN: C, F, I, L;	COMPLEX (MHC PROTEIN/ANTICEN) DRA, DRB1 01010; COMPLEX (MHC PROTEIN/ANTIGEN), HISTOCOMPATIBILITY ANTIGEN
851	1bx2	В	32	147	1.60E-69	-0.22	0.87		HLA-DR2; CHAIN: A, D; HLA- DR2; CHAIN: B, E; HLA-DR2; CHAIN: C, F;	IMMUNE SYSTEM HLA-DK4, MYELIN BASIC PROTEIN, MULTIPLE SCLEROSIS, 2 AUTOIMMUNITY, IMMUNE SYSTEM
851	1bx2	В	32	157	1.60E-69			52.4	HLA-DR2; CHAIN: A, D; HLA- DR2; CHAIN: B, E; HLA-DR2; CHAIN: C, F;	IMMUNE SYSTEM HLA-DK4, MIEDLIN BASIC PROTEIN, MULTIPLE SCLEROSIS, 2 AUTOIMMUNITY, MIMUNE SYSTEM
851	145 E	æ	31	147	4.80E-62	-0.33	0.92		HLA CLASS II HISTOCOMPATIBILITY ANTIGEN; CHAIN: A; HLA CLASS II HISTOCOMPATIBILITY ANTIGEN; CHAIN: B; ENTEROTOXIN TYPE B; CHAIN: C; PEPTIDE INHIBITOR: CHAIN: D;	IMMUNE SYSTEM HLA-DR4, DLA-DR4, DR4; SEB, SUPERANTIGEN; COMPLEX (MHC CLASS IL/SUPERANTIGEN), IMMUNE SYSTEM
851	<u>-</u> [5]	В	30	147	3.20E-67	-0.31	0.82		MAJOR HISTOCOMPATIBILITY	IMMUNE SYSTEM MHC CLASS II DR2A

und PDB annotation	IAIN; FY AIN; ABASIC F:	E IAIN:	- +	AD; CHAIN: MHC II, CLASS II MITC, 1-7, AD; CHAIN: MAC II, CLASS II MITC, 1-7, OVALBUMIN PEPTIDE OVALBUMIN PETIDE OVALBUMIN PEPTIDE OVALBUM	\dashv				-AD; CHAIN: MHC II MHC II, CLASS II MHC I-AD	HAIN: NULL; BLOOD COAGULATION BLOOD COAGULATION, EGF, HYDROLASE,			CHAIN: I, H; PROTEASE, COMPLEX, CO-FACTOR, 2 UE FACTOR; PRECEPTOR ENZYME, INHIBITOR, PRE-PHE-PHE-PHE-PHE-PHE-PHE-PHE-PHE-PHE-PH
Coumpound	COMPLEX ALPHA CHAIN; CHAIN: A, D; MAJOR HISTOCOMPATIBILITY COMPLEX BETA CHAIN; CHAIN: B, E; MYELIN BASIC PROTEN: CHAIN: C, F;	MHC CLASS II I-AK; CHAIN: A, B, P, HEN EGGWHITE LYSOZYME PEPTIDE	MHC CLASS II I-AD; CHAIN: A, B;	MHC CLASS II I-AD; CHAIN: A, B;	MHC CLASS II I-EK; CHAIN: A, B, C, D;	MHC CLASS II I-EK; CHAIN: A, B, C, D;	MHC CLASS II I-EK; CHAIN: A, B. C, D;	MHC CLASS II I-AD; CHAIN: A. B.	MHC CLASS II I-AD; CHAIN: A, B;	FACTOR VII; CHAIN: NULL;	INVASIN; CHAIN: A;	GLYCOSYLTRANSFERASE CYCLODEXTRIN GLUCANOTRANSFERASE (E.C.2.4.1.19) (CGTASE) 1CYG	BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D-PHE-PHE-
SeqFold score			52.61			50.29		52.73					
PMF		0.46		0.71	6.0		0.87		_	0.17	-0.19	-0.19	0.04
Verify		-0.4		-0.39	-0.27		-0.43		-0.5	-0.42	0.05	0.07	-0.24
PSI- BLAST		4.80E-53	9.60E-57	9.60E-57	1.40E-63	1.40E-63	8.00E-64	3.20E-58	3.20E-58	0.0072	3.60E-19	7.20E-15	0.009
End		147	157	147	147	157	147	157	147	474	229	226	475
Start		36	23	35	29	∞	24	22	26	443	2	17	404
Chain ID		В	В	В	В	В	В	В	В		¥		٦
PDB ID		liak	liao	liao	liea	lica	licb	2iad	2iad	1669	lcwv	lcyg	ldan
SEQ ID	Ö	851	851	851	851	851	851	851	851	856	856	856	856

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	2	9	¥ ¥	V	DLASI	200	3			THE TANK A SEGUINDED A SE INHIBITOR
826	ldva	٦	429	472	0.0036	0.07	0.28		DES-GLA FACTOR VIIA (HEAVY CHAIN); CHAIN: H, I; DES-GLA FACTOR VIIA (LIGHT CHAIN); CHAIN: L, M; (DPN)-PHE-ARG; CHAIN: C, D; PEPTIDE E-76; CHAIN:	HYDKOLASEH I DROLASE INIIDIO ON PROTEIN-PEPTIDE COMPLEX
856	leut		18	232	7.20E-13	0.22	-0.19		SIALIDASE; CHAIN: NULL:	HYDROLASE NEURAMINIDASE; HYDROLASE, GLYCOSIDASE
856	1f7e	4	443	475	0.0036	-0.28	0.11		BLOOD COAGULATION FACTOR VII; CHAIN: A;	BLOOD CLOTTING FACTOR VII, BLOOD COAGULATION, EGF-LIKE DOMAIN, BLOOD 2 CLOTTING
856	1hac		429	482	0.0072	-0.21	0		HEREGULIN-ALPHA; CHAIN: NULL;	GROW IH FACTOR NEU DIFFERENTIATION FACTOR (RAT), ACETYLCHOLINE GROWTH FACTOR
856	1hre		230	254	0.0072	-0.7	0.23		GROWTH FACTOR HEREGULIN-ALPHA (EPIDERMAL GROWTH FACTOR-LIKE DOMAIN) 1HRE 3 (NMR, MINIMIZED AVERAGE STRUCTURE) 1HRE 4	
856	1pa m	<	-	171	5.40E-12	0.04	-0.19		CYCLODEXTRIN GLUCANOTRANSFERASE; CHAIN: A, B;	GLYCOSYLTRANSFERASE TRANSFERASE, GLYCOSYLTRANSFERASE, CALCIUM, SIGNAL
857	15u8	<	18	256	3.60E-83	0.85	_		PANCREATIC LIPASE RELATED PROTEIN 2; CHAIN: A;	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
857	1bu8	<	81	276	4.80E-65	0.81			PANCREATIC LIPASE RELATED PROTEIN 2; CHAIN: A;	HYDROLASE HYDROLASE, LIFID DEGRADATION, PANCREATIC LIPASE
857	1cth	∢	8-	276	1.60E-66			167.54	TRIACYLGLYCEROL ACYL- HYDROLASE, CHAIN: A. C; COLIPASE, CHAIN: B, D	COMPLEX (HYDROLASE/CUFACTOR) TRIACYLGLYCEROL LIPASE; COMPLEX (HYDROLASE/COFACTOR). LIPID DEGRADATION
857	let h	∢	88	276	1.60E-66	0.65	-		TRIACYLGLYCEROL ACYL- HYDROLASE, CHAIN: A, C; COLIPASE, CHAIN: B, D	COMPLEX (HYDROLASE/COFACTOR) TRIACYLGLYCEROL LIPASE; COMPLEX (HYDROLASE/COFACTOR), LIPID DEGRADATION

PDB annotation	SERINE ESTERASE RELATED PROTEIN 2 LIPASE; SERINE ESTERASE, HYDROLASE, LIPID DEGRADATION, PANCREAS, 2 GLYCOPROTEIN, CHIMERIC	SERINE ESTERASE RELATED PROTEIN 2 LIPASE; SERINE ESTERASE, HYDROLASE, LIPID DEGRADATION, PANCREAS, 2 GLYCOPROTEIN, CHIMERIC	SERINE ESTERASE RELATED PROTEIN 2 LIPASE; SERINE ESTERASE. HYDROLASE, LIPID DEGRADATION. PANCREAS, 2 GLYCOPROTEIN, CHIMERIC				
Coumpound	RP2 LIPASE; CHAIN: NULL;	RP2 LIPASE, CHAIN: NULL;	RP2 LIPASE, CHAIN: NULL;	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) (TRIACYLGLYCEROL HYDROLASE) 1HPL 3	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED 1LPB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) 1LPB 4	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.13) COMPLEXED WITH COLIPASE AND INHIBITED ILPB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) ILPB 4	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED 11.PB 3 BY
SeqFold score	176.81			167		173.41	
PMF		-	ļ		 		_
Verify		0.87	0.76		0.83		0.81
PSI- BLAST	1.80E-80	1.80E-80	3.20E-65	1.60E-76	1.10E-79	1.10E-79	1.60E-65
End AA	276	256	276	274	256	276	276
Start AA	81	61	61	18	81	81	8.
Chain ID				<	æ	В	æ
PDB ID	lgpi	1gpl	1gpl	Ihpl	11pb	11pb	1lpb
SEQ B NO.	857	857	857	857	857	857	857

SEO	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
2 0 1	9	9	¥Υ	AA.	BLAST	score	score	score		
NO:									UNDECANE PHOSPHONATE METHYL ESTER (TWO	
857	14 <u>1</u>		81	257	5.40E-83	69.0	-		PANCREATIC LIPASE RELATED PROTEIN 1;	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
857	141		18	274	5.40E-83			183.05	CHAIN: NULL; PANCREATIC LIPASE RELATED PROTEIN 1; CHAIN: NULL;	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
857	<u>e</u> .		18	276	1.60E-63	0.71	_		PANCREATIC LIPASE RELATED PROTEIN 1; CHAIN: NULL;	HYDROLASE HYDROLASE, LIPID DEGRADATION, PANCREATIC LIPASE
859	lub1	В	14	156	0.00036	-0.08	0		SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT
859	lerj	V V	151	308	1.30E-44	0.28	0.07		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C.	TRANSCRIPTION INHIBITOR BEIA- PROPELLER
				-			5		CT AT DHA/GLAI PHA	COMPLEX (GTP-
829	l got	æ	132	298	8.00E-46	0.19	0.12		CHIMERA; CHAIN: A; GT-	BINDING/TRANSDUCER) BETAI,
									BETA; CHAIN: B; U1- GAMMA; CHAIN: G;	GAMMAI, TRANSDUCIN GAMMA
										SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN,
										HETEROTRIMER 2 SIGNAL TRANSDUCTION
859	1got	В	∞	309	1.10E-40			52.34	GT-ALPHA/GI-ALPHA	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1,
									BETA: CHAIN: B; GT-	TRANSDUCIN BETA SUBUNIT;
									CAMMIN'S CHAMIN'S C.	SUBUNIT, COMPLEX (GTP-
										HETEROTRIMER 2 SIGNAL TRANSDUCTION
829	lgot	B	95	309	1.10E-40	0.13	-0.11		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT-	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1,
_									BETA; CHAIN: B; GT-	TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA
										SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN,
		_	_							

PDB annotation	HETEROTRIMER 2 SIGNAL TRANSDUCTION	OXIDOREDUCTASE ENZYME, NITRITE REDUCTASE, OXIDOREDUCTASE, DENITRIFICATION, 2 ELECTRON TRANSPORT, PERIPLASMIC	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER. DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA)	FINGER, DNA-BINDING PROTEIN	Cos de par camio parona maia	COMPLEX (ZINC FINGERDINA) COMPLEX (ZINC FINGERDNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEY (7PIC ENICED (DNA) 2INC	FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	CLINC FINGENDINA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2
Coumpound		CYTOCHROME CDI NITRITE REDUCTASE; CHAIN: A, B;	QGSR, ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE PRIDANC SITE: CHAIN: B C:	BINDING SITE, CHAIN. B, C,	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER	OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C.	DINDING SILL, CHAIR, E, C,	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING STTE: CHAIN: B. C.	BINDING SITE, CHAIN. B, C,	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	PROTEIN, CHAIN: C, F, G;
SeqFold score							:											_				
PMF score		-0.12	0.15					86.0			6:00	000	0.0				0.52				_	
Verify score		0	-0.3	0.04		0.48		0.27			0.41		-0.35			,	-0.16				0.1	
PSI- BLAST		1.30E-14	1.60E-25	1.80E-39		9.00E-38		3.60E-39			6.40E-31		1.105-39				3.20E-43				1.40E-44	
End		304	221	277		306		640			64		193				221				249	
Start		221	141	197		225		995			260		7117				140				891	
Chain ID		∀	<	<		V		٧			<		ر.				ပ				U	
PDB ID		1qks	lalh	lalh		lalh		lalh			lath		9 E >	`			Inc v				<u>н</u>	<u> </u>
SEQ ID	2	859	860	860		860		860			098	3,6	98				098				098	

\ 				10.1	DOL	Verify	PMF	SeaFold	Coumpound	PDB annotation
SEQ	10 E	Chain TO	AA	AA	BLAST	score	score	score		
٦										CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
098	1me y	U	961	277	1.60E-46	0.23	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F. G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
860	Ime y	U	224	305	1.60E-47	0.55	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
860	Ime y	0	252	333	1.40E-48	0.49			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGERDINA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDINA)
860	lme y	O	280	361	3.20E-49	0.62	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGERDINA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
860	Jme y	O	308	389	8.00E-50	0.34	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGERDINA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
860	ر ب م	U	336	417	1.60E-50	0.27	-		DNA, CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA) ZINC FINGER/DNA) ZINC
098	y y	U	364	445	1.60E-50	0.42	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGERODAY) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA) ZINC FINGER/DNA) ZINC
098	, Imc	ပ	392	473	4.80E-50	0.16	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (LINC FINGEROUS) ENCE FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX

- т				T					 -
PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA) TRIC	COMPLEA (ZINC FINGENDIA) EINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERDNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEA (ZINC FINGERODIA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERODIA)	COMPLEX (ZINC FINGERODINA) ZINC FINGER, PROTEIN-DNA INTERACTION. PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERODNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A. B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold score							101.87		
PMF		0.98	-	96:0	-			,	-
Verify score		-0.22	0.1	0.15	0.43	69.0		0.71	0.39
PSI- BLAST		7.20E-33	9.60E-50	3.20E-47	3.20E-47	9.60E-50	8.00E-50	8.00E-50	1.60E-49
End		200	501	528	556	584	585	612	640
Start		392	420	448	476	503	503	531	559
Chain ID		O O	O .	O	S	U	U	ပ	၁
PDB ID	+	Imc y	Ime y	Ime y	Ime y	у	lme y	Ime y	lme y
SEQ ID	NO:	860	098	098	860	098	860	860	860

COMPLEX (ZINC FINGENDINA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2
1 IdVIOU DELINE COMINI
CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA
PROTEIN; CHAIN: C, F, G, CRYSTAL STRUCTURE, COMPLEX
TFIIIA; CHAIN: A, D; 5S
TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B. C. E. F.
1
-0.25 0.12
4.80E-31
258
306
O Y
lmc y 1tf6
098

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PDB annotation		REGULATION/DNA) COMPLEX	(TRANSCRIPTION PEGITI ATTON/DNA), RNA	POI YMERASE III. 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	(TRANSCRIPTION	REGULATION/DNA), KNA	POLYMERASE III, 2 TRANSCRUPTION	INITIATION, ZINC FINGEN FROILER	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEXA	(IKANSCKIPTION	RECOLATION/DIAA), KAYA	NITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-FROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION PEGIT ATTON/DNA) YING-YANG 1;	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2
Coumpound		GENE;	CHAIN: B, C, E. F:			CONTRACTOR A D. SC	PIBOSOMAI RNA GENE:	CHAIN: B C E E				TFIIIA: CHAIN: A. D. 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			WAYL CHAIN C. ADENO-	YYI; CHAIN: C, ADENO:	INITIATOR ELEMENT DNA:	CHAIN: A, B;			VVI: CHAIN: C. ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN A B				YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS FO INITIATOR ELEMENT DNA;	CHAIN: A, B;
SeqFold	score																												-						_			
PMF	score						0.64					970	0.0					_						0					5	0.83				_		69.0		
Verify	score						0.11						1.5					900	}					-0.43						رن ان			-			-0.37		
-ISA	BLAST						8.00E-38					36 207 3	6.40E-33					0 KOE-35	200:					9.60E-29						1.30E-30	_					1.80E-34		
End	V V						565					3	298					707	20					221						249						277		
Start	Ψ¥						421	_					449	_				26.7	<u>;</u>					115			_			148						150		
Chain	е						<						∢						<					ပ						၁						O		
aud	2 0						1166						11196					ì	1110					Jubd						1 ubd						lubd		
000) A	Ö					860	}					860						098					860						098						098	}	

SEQ ID	PDB UD	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold	Coumpound	PDB annotation
NO.										FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX CTRANSCRIPTION REGULATION/DNA)
860	lubd	U	0.21	277	3.20E-32	-0.19	0.93		YYI; CHAIN: C: ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B:	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	lubd	O .	194	333	1.80E-48	-0.07	0.96		YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	lubd	ပ	227	333	4.80E-34	0.19	-		YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
098	1ubd	ပ	260	361	3.20E-34	0.27	-		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	lubd	O	306	445	5.40E-52	0.04	0.88		YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	lubd	၁	316	417	3.20E-34	0.2	-		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DIVA;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
Ö									CHAIN: A, B;	INITIATOR ELEMENT, YY I, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	lubd	U	372	473	4.80E-35	-0.14			YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (IRANSCHIPTION) REGULATION/DNA) YING-YANG I; TRANSCRIPTION MITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	106d	U	390	556	3.60E-42	-0.42	0.53		YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	CUMPLEX (TRANSCILL) REQULATIONDNA) YINGYANG 1; TREQULATIONDNA) YINGYANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REQULATION/DNA)
860	lubd	U	400	501	6.40E-35	0	0.78		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (IKANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	lubd	O	456	556	3.20E-31	-0.07	96:0		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
860	1 ubd	U	481	584	1.80E-47	0.23	-		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	REGULATION/DNA) YING-YANG I; REGULATION/DNA) YING-YANG I; INTATOR ELEMENT, YYI. ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
98	1ubd	υ -	484	584	3.20E-36	0.22	-		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;

PDB annotation	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (IKANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)				COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)
Coumpound	INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	SWIS; CHAIN: NULL;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A: DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score			86.51					
PMF		0.98		-	0.72	96.0	-	
Verify		0.43		90.0-	0.15	-0.04	-0.21	-0.1
PSI- BLAST		5.40E-51	5.40E-51	3.20E-34	0.00016	3.20E-31	3.60E-52	5.40E-63
End		640	641	640	642	276	307	363
Start		529	531	539	615	140	170	197
Chain ID		U	U	U		4	<	<
PDB ID		1ubd	lubd	lubd	1zfd	2gli	2gli	2gli
SEQ ID	SO.	098	098	098	860	860	860	098

					104	Wante.	DMG	Confiold	Commound	PDB annotation
SEQ	20 GI	Chain El Ein	AA	AA	BLAST	score	score	score		
860 860	2gli	<	224	391	3.60E-64	90.0			ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
098	2gli	<	227	360	1.60E-33	0.13	0.98		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
9860	2gli	V V	280	447	1.80E-64	0.25	66:0		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
098	2gli	⋖	308	558	3.60E-57	-0.4	0.19		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
860	2gli	<	316	444	1.30E-34	0.03	0.99		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
860	2gli	<	400	527	1.60E-32	-0.21	0.49		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
098	2gli	4	421	614	5.40E-60	-0.07	0.19		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
860	2gli	∢	456	586	1.30E-30	-0.02	0.99		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
860	2gli	4	484	119	1.40E-35	0.27		·	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D:	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
860	2gli	4	503	640	3.60E-64	0.53	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
860	2gli	٧	503	642	3.60E-64			91.33	ZINC FINGER PROTEIN GLII;	COMPLEX (DNA-BINDING

	- ,	<u>'i</u>	₹.)E		3; G	2; ;	5 <u>9</u>	·z'
notation	E-FINGER GLI; GI PLEX (DNA- DNA)	NDING E-FINGER GLI; GI PLEX (DNA- TDNA)	YME DLH; YDROLASE, OCARBON SRINE ESTERASE, LENEBUTENOLID; ENZYME	EF-TS; TOR, NUCLEOTII 3INDING, 2	LONGATION ATION FACTOR (EAT UNSTABLE, TOR FOR STABLE, TOR, COMPLEX N FACTORS)	IAAB 20	AGA DNA-BINDIN IN A OF RAT HMC (IAAB 20	DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMG1; 1AAB 8 HMG-BOX 1AAB 20	DNA BENDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION,
PDB annotation	PROTEINDNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEINDNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	HYDROLYTIC ENZYME DLH; DIENELACTONE HYDROLASE, AROMATIC HYDROCARBON CATABOLISM, 2 SERINE ESTERASE, CARBOXYMETHYLENEBUTENOLIDA SE, 3 HYDROLYTIC ENZYME	COMPLEX OF TWO ELONGATION FACTORS EF-TU; EF-TS; ELONGATION FACTOR, NUCLEOTIDE EXCHANGE, GTP-BINDING, 2 COMPLET FOR TWO FLONGATION	FACTORS COMPLEX (TWO ELONGATION FACTORS) ELONGATION FACTOR FOR TRANSFER, HEAT UNSTABLE, ELONGATION FACTOR FOR TRANSFER, HEAT STABLE, ELONGATION FACTOR, COMPLEX COMPLEX	DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMGI; 1AAB 8 HMG-BOX 1AAB 20	DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMG1; IAAB 8 HMG-BOX IAAB 20	DNA-BINDING HMGA DNA-BINDING HMG-BOX DOMAIN A OF RAT HMG1 IAAB 8 HMG-BOX IAAB 20	DNA BINDING PR DNA BENDING, D
Coumpound	CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	DIENELACTONE HYDROLASE; CHAIN: NULL;	ELONGATION FACTOR TU; CHAIN: A, B, E, F; ELONGATION FACTOR TS: CHAIN: C, D, G, H;	ELONGATION FACTOR TU; CHAIN: A, C; ELONGATION FACTOR TS; CHAIN: B, D;	HIGH MOBILITY GROUP PROTEIN; IAAB 5 CHAIN: NULL; IAAB 6	HIGH MOBILITY GROUP PROTEIN; 1AAB 5 CHAIN: NULL: 1AAB 6	HIGH MOBILITY GROUP PROTEIN; 1AAB 5 CHAIN: NULL; 1AAB 6	NON HISTONE PROTEIN 6 A; CHAIN: A;
SeqFold									
PMF score		0.46	0.96	0.66	0.75	0.89	_	0.33	0.94
Verify score		-0.16	0.02	0.02	-0.56	-0.09	99.0	0.41	-0.12
PSI- BLAST		1.40E-36	0.00013	0.00018	0.00018	9.00E-18	5.40E-16	1.80E-06	1.80E-16
End		279	338	2676	2673	185	433	290	180
Start		95	255	2646	2646	106	363	528	112
Chain ID		4		Jo					₹.
PDB ID		2gli	1din	laip	lefu	laab	1 aab	laab	1cg7
SEQ 1D	Ö	098	862	864	864	865	865	865	865

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PROTEIN	DNA BINDING PROTEIN HMG BUX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	DNA BINDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	DNA BINDING PROTEIN HWG BCO., DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2 BINDING PROTEIN	DNA BINDING PROTEIN THAN DOOR DNA BENDING, DNA RECOGNITION CHROMATIN, NMR, DNA 2 BINDING PROTEIN	GENE REGULATIONDNA HMG-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	GENE KEGULA HOWDNA HWG-1, AMPHOTERN, HEPARIN-BINDING PROTEIN P30, HIGH-MOBILLITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	GENE KEGULATION/DIA AMPI-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	GENE REGULATION DA DANG-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30, HIGH-MOBILITY GROUP DOMAIN, BENT DNA. PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA
	NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	HIGH MOBILITY GROUP 1 PROTEIN: CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;	HIGH MOBILLITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;	HIGH MOBILLITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(IDO) CHAIN: B; DNA (5'- CHAIN: C;
	0.83	_	0.42	0.24	0.92	0.71	66:0	0.11
	0.3	0.45	-0.4	0.28	-0.29	0.02	0.4	0.03
	3.60E-07	5.40E-16	7.20E-06	1.80E-06	5.40E-14	3.60E-07	3.60E-15	7.20E-05
1	320	433	916	601	177	320	433	510
	262	369	476	532	112	262	369	476
		4	A	-V	<	4	∀	4
	1cg7 A	1cg7	1cg7	1cg7	1ckt	ा ट्स	lckt	lckt
	865	865	865	865	865	865	865	865
		Lcg7 A 262 3.60E-07 0.3 0.83 NON HISTONE PROTEIN 6 A; CHAIN: A; CHAIN: A;	Leg7 A 262 320 3.60E-07 0.3 0.83 NON HISTONE PROTEIN 6 A; Leg7 A 369 433 5.40E-16 0.45 1 NON HISTONE PROTEIN 6 A;	Leg7 A 262 320 3.60E-07 0.3 0.83 NON HISTONE PROTEIN 6 A; CHAIN: A; Leg7 A 369 433 5.40E-16 0.45 1 NON HISTONE PROTEIN 6 A; Leg7 A 476 516 7.20E-06 -0.4 0.42 NON HISTONE PROTEIN 6 A; Leg7 A 476 516 7.20E-06 -0.4 0.42 CHAIN: A;	1cg7 A 262 320 3.60E-07 0.3 0.83 NON HISTONE PROTEIN 6 A; 1 1cg7 A 369 433 5.40E-16 0.45 1 NON HISTONE PROTEIN 6 A; 1 1cg7 A 476 516 7.20E-06 -0.4 0.42 CHAIN: A; CH	1cg7 A 262 320 3.60E-07 0.3 0.83 DIA HISTONE PROTEIN 6 A; 1 CHAIN: A; CHAIN: A; CHAIN: A; CHAIN: A; CHAIN: A; CHAIN: A; DNA (5*- CHAIN: B; DNA (5*- CHAIN: B; DNA (5*- CHAIN: C; DNA (5*- CHAIN: B; DNA (5*- CHAIN: C; DNA (5*-	Leg7 A 262 320 3.60E-07 0.3 0.83 NON HISTONE PROTEIN 6 A; CHAIN: B; CHAIN: A; CHAIN: B; CHAIN: B; CHAIN: B; CHAIN: B; CHAIN: CHAIN: B; CHAIN: C,	1eg7 A 262 320 3.60E-07 0.3 0.83 DON HISTONE PROTEIN 6 A; 1 1 1 1 1 1 1 1 1

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PDB annotation	GENE REGULATION/DNA HMG-1, AMPHOTERIN, HEPARIN-BINDING PROTEIN P30; HIGH-MOBILITY GROUP DOMAIN, BENT DNA, PROTEIN-DRUG-DNA 2 COMPLEX, GENE REGULATION/DNA	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN				COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA)	
Coumpound	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5'- D(*CP*CP*(JDO) CHAIN: B; DNA (5'- CHAIN: C;	ALPHA SPECTRIN; CHAIN: A, B, C;	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING IHME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) IHME 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING 1HME 3 HMG-BOX DOMAIN B OF RAT HMGI) (NMR, 1 STRUCTURE) 1HME 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING 1HME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) 1HME 4	HUMAN SRY; 1HRY 6 CHAIN: A; 1HRY 7 DNA; 1HRY 9 CHAIN: B; 1HRY 10	HUMAN SRY; 1HRY 6 CHAIN: A; 1HRY 7 DNA; 1HRY 9 CHAIN: B; 1HRY 10	DNA-BINDING HIGH MOBILITY GROUP PROTEIN 1 (HMG1) BOX 2,
SeqFold score								
PMF	0.01	0	0.57	0.43	_	0.39	0.78	0.21
Verify score	-0.01	-0.16	0	-0.19	0.04	-0.52	0.19	0.44
PSI- BLAST	1.40E-05	7.20E-08	3.60E-14	0.00036	3.60E-13	9.00E-15	1.80E-16	9.00E-12
End	290	585	091	303	418	180	433	160
Start	530	382	107	257	369	0=	368	112
Chain ID	4	<				<	V	
PDB ID	lckt	Icun	o than	o lhm	o lhm	1hry	lhry	1 hsm
SEQ 15 NO.	865	865	865	865	865	865	865	865

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			GENE REGULATION/DNA HMG-D; PROTEIN-DNA COMPLEX, HMG DOMAIN, NON-SEQUENCE SPECIFIC 2 CHROMOSOMAL PROTEIN HMG-D	GENE REGULATION/DNA HMG-D; PROTEIN-DNA COMPLEX, HMG DOMAIN, NON-SEQUENCE SPECIFIC 2 CHROMOSOMAL PROTEIN HMG-D	GENE REGULATION/DNA LEF-1 HMG; LEF1, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	GENE REGULATION/DNA LEF-1 HMG; LEF1, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	GENE REGULATION/DNA LEF-1 HMG;
COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN I (HMG1) BOX 2, COMPLEXED WITH IHSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	DNA-BINDING HIGH MOBILITY GROUP PROTEIN I (HMG1) BOX 2, COMPLEXED WITH IHSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	DNA (5'- D(*GP*CP*GP*AP*TP*AP*TP *CP*GP*C)-3'); CHAIN: C, D; HIGH MOBILITY GROUP PROTEIN D; CHAIN: A, B;	DNA (5'- D(*GP*CP*GP*AP*TP*AP*TP *CP*GP*C)-3'); CHAIN: C, D; HIGH MOBILITY GROUP PROTEIN D: CHAIN: A, B;	LYMPHOID ENHANCER- BINDING FACTOR; CHAIN: A; DNA (5'- CHAIN: B; DNA (5'- CHAIN: C;	LYMPHOID ENHANCER- BINDING FACTOR; CHAIN: A; DNA (5'- CHAIN: B; DNA (5'- CHAIN: C;	LYMPHOID ENHANCER-
	0.37		0.01	0.78	0.36	0.58	1
	0.02	9.0	-0.69	0.18	-0.23	11.0	0.76
	0.00036	5.40E-13	1.10E-10	5.40E-12	1.80E-14	3.60E-07	1.10E-14
	303	81	153	411	180	320	433
	262	369	112	367	113	262	371
			<	4	V V	₹	٧
	1hsm	Ihsm	lqrv	Iqrv	21cf	2lef	2lcf
2	865	865	865	865	865	865	865
	COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	1hsm 262 303 0.00036 0.02 0.37	Thism 262 303 0.00036 0.02 0.37 DIA-BINDING HIGH	Ihism 262 303 0.00036 0.02 0.37 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE	Ihism 262 303 0.00036 0.02 0.37 MGRACAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) IHSM 4	Ihsm 262 303 0.00036 0.02 0.37 DNA-BINDING HIGH	Ihsm 262 303 0.00036 0.02 0.37 MINIMIZED AVERAGE

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PDB annotation	LEFI, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	GENE REGULATION/DNA LEF-1 HMG; LEF1, HMG, TCR-A, TRANSCRIPTION FACTOR, DNA BINDING, DNA 2 BENDING, COMPLEX (HMG DOMAIN/DNA), GENE REGULATION/DNA	COMPLEX (TRANSDUCERTRANSDUCTION) GT BETA-GAMMA; MEKA, PP33; PHOSDUCIN, TRANSDUCIN, BETA-GAMMA, SIGNAL TRANSDUCTION, 2 REGULATION, PHOSPHORYLATION, G PROTEINS, THIOREDOXIN, 3 VISION, MEKA, COMPLEX (TRANSDUCER/TRANSDUCTION)		ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	ISOMERASE ISOMERASE, MUTASE, INTRAMOLECULAR TRANSFERASE	ISOMERASE ISOMEKASE, MULASE, INTRAMOLECULAR TRANSFERASE	TRANSCRIPTION REGULATION SIGMATO; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION		TRANSFERASE GLYCOSYLTRANSFERASE
Coumpound	BINDING FACTOR; CHAIN: A; DNA (5'-CHAIN: B; DNA (5'-CHAIN: C;	LYMPHOID ENHANCER- BINDING FACTOR; CHAIN: A; DNA (5'- CHAIN: B; DNA (5'- CHAIN: C;	TRANSDUCIN; CHAIN: B, G; PHOSDUCIN; CHAIN: P;		SYNTAXIN-1A; CHAIN: A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	METHYLMALONYL-COA MUTASE; CHAIN: A, B, C, D;	MUTASE; CHAIN: A, B, C, D;	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;		SPORE COAT POLYSACCHARIDE BIOSYNTHESIS PROTEIN CHAIN: A;
SeqFold score											
PMF score		0.68	-0.19		-0.2	-0.19	-0.2	-0.19	-0.19		0.22
Verify score		0.28	0.78		0.53	0.58	0.36	0.64	0.09	_	0.06
PSI- BLAST		3.60E-06	3.60E-12		1.80E-09	5.40E-09	3.60E-12	1.80E-10	1.80E-08		0.0018
End		290	709		465	467	464	482	463	_	302
Start		532	639		396	396	390	396	386		97
Chain ID		*	م.		<	₹	∢_	<			∢
PDB ID		2lef	2trc		lez3	lquu	Ireq	lreq	1sig		1989
SEQ	Ö	865	865		866	998	866	998	998		871

	 1				 -j	_		
PDB annotation	RNA-BINDING PROTEIN/RNA TRA PRE-MRNA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	GENE REGULATION/RNA POLY(A) BINDING PROTEIN I, PABP I; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA	RNA BINDING PROTEIN RNA- BINDING DOMAIN	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1, NUCLEAR PROTEIN, HNRNP, RBD, RRM, RNP, RNA BINDING, 2 RIBONUCLEOPROTEIN	
Соитроипа	SXL-LETHAL PROTEIN; CHAIN: A, B; RNA (5'- R(P*GP*UP*UP*UP*U P*UP*UP*UP*UP*U)- CHAIN: P, Q;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D. E. F. G. H: RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A,3'); CHAIN: M, N, O, P, Q, R, S, T;	POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	HU ANTIGEN C; CHAIN: A;	HNRNP A1; CHAIN: NULL;	RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND
SeqFold score								
PMF	-	0.92	-		-	1.	_	0.92
Verify score	0.52	0.52	0.99	0.92	0.95	0.83	0.48	0.56
PSI- BLAST	1.40E-21	6.40E-24	1.60E-20	1.60E-20	1.60E-20	1.10E-21	3.20E-22	1.40E-21
End	104	110	106	106	106	901	124	109
Start AA	9	9	27	27	27	22	21	24
Chain ID	¥.	V	В	(L.	±	Ą		
PDB ID	1b7f	Icvj	Icvj	lcvj	lcvj	148z	lhal	lsxl
SEQ ID NO:	872	872	872	872	872	872	872	872

PDB annotation		COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	RNA BINDING DOMAIN RNA BINDING DOMAIN, RBD, RNA RECOGNITION MOTIF, RRM, 2 SPLCING INHIBITOR, TRANSLATIONAL INHIBITOR, SEX 3 DETERMINATION, X CHROMOSOME DOSAGE COMPENSATION	CALCIUM/PHOSPHOLIPID BINDING PROTEIN PI1, CALPACTIN LIGHT CHAIN; S100 FAMILY, EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2 CALCIUM/PHOSPHOLIPID BINDING PROTEIN	CALCIUM-BINDING CNTNC; CALCIUM-BINDING, REGULATION, TROPONIN C, CARDIAC MUSCLE 2 CONTRACTION	CALCIUM BINDING CALCIUM BINDING	METAL BINDING PROTEIN CAVP; EF- HAND FAMILY, CALCIUM BINDING PROTEIN, NMR		
Coumpound	RNA-BINDING DOMAIN 1SXL 3 (RBD-2), RESIDUES 199 - 294 PLUS N-TERMINAL MET) 1SXL 4 (NMR, 17 STRUCTURES) 1SXL 5	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	SEX-LETHAL; CHAIN: A, B, C;	S100A10; CHAIN: A, B;	CARDIAC N-TROPONIN C; CHAIN: NULL;	CALCIUM-BINDING PROTEIN; CHAIN: NULL;	CALCIUM VECTOR PROTEIN; CHAIN: A;	CALCIUM-BINDING PROTEIN CALBINDIN D9K (INTACT FORM) (NMR, 13 STRUCTURES) 1CBI 3	CALCIUM-BINDING PROTEIN CALMODULIN (VERTEBRATE) 1CLL 3
SeqFold score									
PMF score		0.99	66.0	-	0.29	0.15	0.16		0.16
Verify score		0.68	0.58	0.66	-0.25	-0.05	-0.22	0.35	-0.14
PSI- BLAST		1.30E-22	1.60E-20	7.20E-18	9.00E-07	5.40E-07	7.20E-07	9.00E-22	3.60E-07
End		124	104	93	77	78	78	83	92
Start		61	9	=	21	22	22	8	=
Chain TD		V	A	<			¥		
PDB ID		2up1	3sxl	la4p	lap4	1bu3	1c7w	1cb1	1cli
SEQ NO.		872	872	873	873	873	873	873	873

								 -	
PDB annotation	METAL BINDING PROTEIN YEAST FREQUENIN EF-HAND, CALCIUM	METAL TRANSPORT MRPS, S100AS, CALGRANULIN A CALCIUM-BINDING PROTEIN, CRYSTAL STRUCTURE, MAD, MIGRATION 2 INHIBITORY FACTOR_RELATED PROTEIN 8, S100 PROTEIN	COMPLEX (LIGAND/ANNEXIN) CALGIZZARIN, S100 FAMILY, EF- HAND PROTEIN, COMPLEX (LIGAND/ANNEXIN), 2 LIGAND OF ANNEXIN II, CALCIUM/PHOSPHOLIPID BINDING PROTEIN	HYDROLASE CEREBROSIDE-3- SULFATE-SULFATASE; CEREBROSIDE-3-SULFATE HYDROLYSIS, LYSOSOMAL ENZYME, 2 HYDROLASE	LYASE DELTA3,5,DELTA2,4- DIENOYL-COENZYME A ISOMERASE, LYASE, DIENOYL-COA ISOMERASE	LYASE DELTA3,5,DELTA2,4- DIENOYL-COENZYME A ISOMERASE, LYASE, DIENOYL-COA ISOMERASE	LYASE METHYLMALONYL COA, DECARBOXYLASE	LYASE DEHALOGENASE; LYASE	LYÁSE CROTONASE, ENOYL-COA HYDRATASE I; LYASE, HYDRATASE, B-OXIDATION, FATTY ACID DEGRADATION, COA, 2 LIGAND BINDING
Coumpound	CALCTUM-BINDING PROTEIN NCS-1; CHAIN: A;	MIGRATION INHIBITORY FACTOR-RELATED PROTEIN 8; CHAIN: A, B;	S100C PROTEIN; CHAIN: A; ANNEXIN I; CHAIN: D;	ARYLSULFATASE A; CHAIN: NULL;	DIENOYL-COA ISOMERASE; CHAIN: A, B, C;	DIENOYL-COA ISOMERASE; CHAIN: A, B, C;	METHYLMALONYL COA DECARBOXYLASE; CHAIN: A, B, C;	4-CHLOROBENZOYL COENZYME A DEHALOGENASE; CHAIN: A, B. C.	2-ENOYL-COA HYDRATASE; CHAIN: A. B. C, D. E, F.
SeqFold score		1							
PMF score	0.01	0.81	_	0.46	0.17	0.35	0.01	-0.11	0.13
Verify	-0.07	0.15	0.38	0.29	-0.17	-0.02	-0.23	0.3	-0.1
PSI- BLAST	9.00E-08	1.80E-17	1.80E-18	9.60E-65	1.80E-14	3.20E-22	1.60E-18	4.80E-21	3.20E-24
End	81	93	55	330	121	144	145	144	144
Start AA	=	_	∞	39	48	55	45	47	47
Chain ID	V	<	4		<	<	₹	¥.	A
PDB ID	1fpw	1mr8	lqís	lauk	1dci	1dci	lcf8	lnzy	2dub
SEQ ID	873 373	873	873	874	875	875	875	875	875

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uo	ON UBC2; ON, NG ENZYN	ON UBC2; ON, NG ENZYN	QUITIN-	QUITIN-	S, YEAST	0 4 01.	COMPLEX	SE, PROTE PR MOTIF		DOMAIN, LICAL	EIN BINDU	DOMAIN,	LICAL 9, PROTEIN	PROXISMO	EROXIN-5	COMPLE)	ŒPEAT, TF	EROXISMC	PEROXIN-5	COMPLEY	REPEAT, 11			OPANE	INONE TO
PDB annotation	NJUGATI NJUGATI NJUGATI	NJUGATI NJUGATI NJUGATI	UITIN, UB G ENZYMI	UITIN, UB	G ENZYMI	VO TO VO	NALLING	PHOXIDA		HOP, TPR- IPLEX, HE	90, 2 PROT	HOP, TPR-	л., 2 нsр7.	ROTEIN P	PTS1-BP, 1	N-PEPTIDE	PEPTIDE I 'F A T	ROTEIN P.	PTS1-BP,	N-PEPTIDI	PEPTIDE I	101	CTASE	CTASE, TR	OF 2 TROF
Id	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	0.61	SIGNALLING COMPLEX KACT; P67PHOX; SIGNALLING COMPLEX,	GTPASE, NADPH OXIDASE, PROTEIN- PROTEIN 2 COMPLEX TPR MOTIF		CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL	REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN,	PEPTIDE-COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN BINDING	SIGNAL ING PROTEIN PEROXISMORE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	PTS1 PROTEIN-PEPTIDE COMPLEX,	TETRATRICOPEPTIDE REPEAT, TPR, 2 HEI ICAL REPEAT	SIGNALING PROTEIN PEROXISMORE	RECEPTOR 1, PTS1-BP, PEROXIN-5,	PTS1 PROTEIN-PEPTIDE COMPLEX	TETRATRICOPEPTIDE REPEAT, 1PK, BELICAL BEPFAT	וברוכער עבו	OXIDOREDUCTASE	OXIDOREDUCTASE, TROPANE	REDUCTION OF 2 TROPINONE TO TROPINE, SHORT-CHAIN
							.⊼ 	<u>ة</u> ك				S	<u>~</u> ≪ ¤	+	_	<u>a</u>		╁		<u></u>		-	 		
pu	JGATING HAIN: A,	JGATINC HAIN: A,	JGATINC A;	JGATINC A;	JGATINC A;		Z	IAIN: A;	CHAIN:	OF HOP; PEPTIDE	3;	HOP;	70. C, D;	APGETTA	FOR:		TIDE;	ARGETIN	TOR:	<u>.</u>	TIDE;		UCTASE.		
Coumpound	N-CONJ(RAD6; C	N-CONIC RAD6; C	N CONJI CHAIN:	CHAIN	CHAIN	0000	ATED C3 UM TOX	TE 1; CF	2 (NCF-2)	OMAIN (CHAIN: 1	MAIN OF	CHAIN:	CMAN T	RECEPT	, B; PTS	VING PER	OMAL T	RECEP	V. B; PTS	ZING PEI	, 0,	NE RED	γ, Β,	
9	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;		RAS-RELATED C3 BOTULINUM TOXIN	SUBSTRATE 1; CHAIN: A;	FACTOR 2 (NCF-2) CHAIN: B;	TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE	MEEVD; CHAIN: B;	TPRI-DOMAIN OF HOP;	CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;	BEBOYISOMAI TABGETING	SIGNAL I RECEPTOR:	CHAIN: A, B; PTS1-	CONTAINING PEPTIDE;	PEROXISOMAL TARGETING	SIGNAL I RECEPTOR:	CHAIN: A. B; PTSI-	CONTAINING PEPTIDE;	CHAIN	TROPINONE REDUCTASE-I;	CHAIN: A, B;	
SeqFold score	139.32		226.34																						
PMF		-		_	-	Ī	-0.1 4		_	-0.15		-0.19		0.5	-0.18			6. 8.	:				_		
Verify		0.84		16.0	0.71		0.15			0.31		0.02		,	40.0			0.01					0.35		
PSI- BLAST	3.60E-63	3.60E-63	9.00E-63	9.00E-63	4.80E-62		3.20E-09			1.60E-18		8.00E-09		1, 000	4.80E-14			1 40F-23	7-701-1				6.40E-29		
End	151	146	147	147	146		137			112		153) cr			241					147		
Start	-	2	-	2	3		12			7		54			-			12	71		_		33		
Chain ID	<	<	<	<	∢		æ		_	٧		A			۷			4	۲				A		
PDB TD	layz	layz	1qcq	lqcq	lqcq		1.00 F.+96	 : :		le!r		lelw			Itch	_		1 fr.h	<u> </u>				lac		
SEQ D	877	877	877	877	877		878			878		878		9	8/8			878	0				881		

Coumpound FDB annotation	Н	TROPINONE REDUCTASE-1; OXIDOREDUCTASE CHAIN: A, B; ALKALOID BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO TROPINE, SHORT-CHAIN DEHYDROGENASE	GENASE; CHAIN:	HENYL-2,3- RODIOL-2,3- ROGENASE; CHAIN:	CTASE;	7 ALPHA- OXIDOREDUCTASE SHORT-CHAIN HYDROXYSTEROID DEHYDROGENASE, BLE ACID CATABOLISM A, B;	3- IHDC 4	SEPIAPTERIN REDUCTASE; OXIDOREDUCTASE SEPIAPTERIN CHAIN: NULL; REDUCTASE, TETRAHYDROBIOPTERIN,		TRIHYDROXYNAPHTHALEN OXIDOREDUCTASE E REDUCTASE; CHAIN: A, B; REDUCTASE; OXIDOREDUCTASE
TROPINONE RED CHAIN: A, B; ALCOHOL DEHYDROGENA A, B;	TROPINONE RED CHAIN: A, B; ALCOHOL DEHYDROGENA A, B;	ALCOHOL DEHYDROGENA A. B;		CIS-BIPHENYL-2,3- DIHYDRODIOL-2,3- DEHYDROGENASE; NULL;	CARBONYL REDU CHAIN: A, B, C, D;	7 ALPHA- HYDROXYSTEROID DEHYDROGENASE; A, B;	OXIDOREDUCTASE ALPHA, 20-BETA- HYDROXYSTEROID DFHYDROGENASE (E.C.I.1.53) IHDC 3 COMPLEXED WITH CARBENOXOLONE	SEPIAPTERIN R CHAIN: NULL;	E REDUCTASE;	TROPINONE REDUCTASE-II:
			•							
6.0				32 0.93	1 0.98	0.68	0.34 0.99	0.47 0.9	0.38	0.49 0.99
29 0.36				24 0.32	21 0.1				<u> </u>	Γ
6.40E-29	6.40E-2	1.40E-2		1.60E-24	4.80E-21	1.60E-34	1.10E-29	1.80E-20	4.80E-33	6.40E-29
		147	129	147	148	147	148	129	147	147
33	33		38	36	36	31	35	38	23]=
		В	\		4	4	«		<	4
		lacı	1616	1bdb	1cyd	Ifmc	Ihdo	loaa	lybv	28e2
	2	188	881	881	188	881	188	881	881	188

SEQ D	PDB ID	Chain ID	Start	End AA	PSI- BLAST	Verify score	PMF	SeqFold	Coumpound	PDB annotation
Ö									CHAIN: A, B;	OXIDOREDUCTASE, TROPANE ALKALOID BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO PSEUDOTROPINE, SHORT-CHAIN DEHYDROGENASE
										COULTINOT CONTEST OF THE CONTEST OF
882	layz	∢	9	157	1.10E-45	0.49	_		UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B,	UBIQUITIN CONJUGATION UBCZ; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
882	layz	A	9	171	1.10E-45			90.32	UBIQUITIN-CONJUGATING	UBIQUITIN CONJUGATION UBC2:
									ENZYME KADO; CHAIN: A, B, C;	UBIQUITIN-CONJUGATING ENZYME
882	1042	Ω	10	157	6.40E-39	0.3	_		UBIQUITIN-PROTEIN	LIGASE E6AP; UBCH7; BILOBAL
<u></u>	!	ı							LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING	STRUCTURE, ELONGATED SHAPE, ES UBIQUITIN LIGASE, E2 2 UBIQUITIN
									ENZYME E2; CHAIN: D;	CONJUGATING ENZYME
882	1c4z	Ω	10	166	6.40E-39			89.72	UBIQUITIN-PROTEIN	LIGASE E6AP; UBCH7; BILOBAL STRIICTIIRE E1 ONGATED SHAPE. E3
									UBIQUITIN CONJUGATING	UBIQUITIN LIGASE, E2 2 UBIQUITIN
									ENZYME E2; CHAIN: D;	CONJUGATING ENZYME
882	lqcq	4	7	168	9.60E-49			87.99	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
882	Iqcq	¥	∞	157	9.60E-49	9.0	-		UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
882	lu9a	<	3	168	9.60E-43			74.26	UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME
										UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME;
										UBIQUITIN-DIRECTED 2
										PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
882	lu9a	A	5	157	9.60E-43	0.17	-		UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME
										UBIQUITIN-CONJUGATING ENZYME;
										UBIQUITIN-DIRECTED 2
										PROTEOLYSIS, CELL CYCLE CONTROL: LIGASE
882	2aak		~	157	6.40E-48	0.38	_		UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION UBCI;
882	2aak		9	148	6.40E-48		_	93.12	UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION UBCI:
6			-	731	2 JOE 44			91 53	INCYME; CHAIN: NOLL;	UBIQUITIN CONTIGATION
788	76.20		-	2	3.20E-44			61.33	Opinion in Colors	

PDB	3 Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold	Coumpound	PDB annotation
								ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
	2c2c	3	157	3.20E-44	0.27	-		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
2ncz		9	157	3.20E-51	0.63	_		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
2ucz		7	165	3.20E-51			121.4	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
layz	Y Z	9	178	6.40E-50			114.6	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C,	UBIQUITIN CONJUGATION UBC2: UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
layz	4	9	178	6.40E-50	0.73	_		UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION UBIQUITIN-CONJUGATING ENZYME
1042	Q 2	01	165	3.20E-42	0.4			UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
1042	Z D	10	691	3.20E-42			104.56	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONTUGATING ENZYME
l dcd	۷ b	7	166	1.10E-53			104.4	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN. CONJUGATING ENZYME, YEAST
1 dcd	A P	∞	166	1.10E-53	0.53	-		UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
1u9a	K	m.	174	4.80E-45			94.76	ÜBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
6	lu9a A	8	173	4.80E-45	0.8	-		UBC9, CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ВŞ	e	2	AA	Ϋ́	BLAST	score	score	score		
										PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
882	2aak		5	172	8.00E-52	0.62	-		UBIQUITIN CONJUGATING ENZYME: CHAIN: NULL:	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
882	2aak		9	174	8.00E-52			116.78	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
882	2c2c		-	176	1.60E-45			109.63	UBIQUITIN CONJUGATING	UBIQUITIN CONTUGATION
									ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION,
										UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
882	2e2c		3	160	1.60E-45	0.55	_		UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION
-									ENZYME: CHAIN: NULL;	UBIQUITIN CONJUGATION.
										UBIQUI IN CARRIER PROTEIN. THIOESTER 2 BOND, LIGASE
882	2ucz		9	160	1.60E-55	0.59	-		UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION UBC7,
									ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION, LIGASE, YEAST
882	2ucz		7	174	1.60E-55			148.79	UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION UBC7;
									ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION, LIGASE,
882	lavz	A	9	128	4.80E-37	0.19	0.75		UBIOUITIN-CONJUGATING	UBIQUITIN CONJUGATION UBC2;
	,	!							ENZYME RAD6; CHAIN: A, B,	UBIQUITIN CONJUGATION,
									C;	UBIQUITIN-CONJUGATING ENZYME
882	1c4z	Ω	01	128	1.60E-28	-0.22	0.54		UBIQUITIN-PROTEIN	LIGASE E6AP; UBCH7; BILOBAL
									LIGASE E3A; CHAIN: A, B, C;	STRUCTURE, ELONGATED SHAPE, E3
							_		UBIQUITIN CONJUGATING	UBIQUITIN LIGASE, E2 2 UBIQUITIN
000		٤	٤		00 40%				ENZIME EZ; CHAIN: U;	TICAGE FLAB. 110CHT. BILOBAL
788	lc4z	<u>a</u>	2		1.60E-28			60.16	UBIQUITIN-PROTEIN	LIGASE ESAF, UBCHI, BILUBAL
						_			UBIQUITIN CONJUGATING	UBIQUITIN LIGASE, E2 2 UBIQUITIN
									ENZYME E2; CHAIN: D;	CONJUGATING ENZYME
882	1dcd	٧	∞	128	3.20E-39	0.2	9.0		UBIQUITIN CONJUGATING	LIGASE UBIQUITIN, UBIQUITIN-
000	-		,	955	1 200 24	9.0	5		THE CHAIN A	THEOTHER CONTINGATING ENTANCE
288	I uya	<	^	871	3.20E-34	<u>နှ</u>	0.72		UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENCYME
										TIBIOTITIN-CONTIGATING ENZYME
								_		UBIQUITIN-DIRECTED 2
		-··								PROTEOLYSIS, CELL CYCLE
882	2aak		5	128	3.20E-38	90:0	0.55		UBIQUITIN CONJUGATING	UBIQUITIN CONJUGATION UBCI.
									ENZYME: CHAIN: NULL;	UBIQUITIN CONJUGATION, LIGASE

PDB annotation	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7: UBIQUITIN CONJUGATION, LIGASE, YEAST	The state of the s	UBIQUITIN CONJUGATION UBCZ; UBIQUITIN CONJUGATION UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION UBIQUITIN-CONJUGATING ENZYME	LIGASE E6AP, UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME	TYOAGE POAD, TROUTS, DIT OBAI	LIGANE EBAP', UBCH', BILUBALL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONIUGATING ENZYME	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2
Coumpound	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	ÜBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;		UBIQUITIN-CONJUGATING ENZYME RAD6, CHAIN: A, B, C;	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONIUGATING ENZYME F2: CHAIN: D:	Citation De, Citation D.	UBIQUITIN-PROTEIN LIGASE E3A, CHAIN: A, B, C, UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	UBC9; CHAIN: NULL;	UBC9, CHAIN: NULL:
SeqFold score	50.62			56.09			90.32			89.72	87.99		74.26	
PMF		0.25	0.49			-						-		_
Verify score		0.17	-0.09			0.49		0.3				9.0		0.17
PSI- BLAST	3.20E-38	1.10E-35	1.60E-35	1.60E-35		1.10E-45	1.10E-45	6.40E-39		6.40E-39	9.60E-49	9.60E-49	9.60E-43	9.60E-43
End	155	126	128	136		157	171	157		166	168	157	168	157
Start	9	3	9	7		9	9	10		01	7	8	E.	\$
Chain ID						₹	<	۵		Ω	4	<	<	A
PDB ID	2aak	2e2c	2ucz	2ucz		layz	layz	1c4z		1c4z	lqcq	ldcd	1u9a	lu9a
SEQ	882	882	882	882		883	883	883		883	883	883	883	883

PDB annotation	PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE	UBIQUITIN CONJUGATION	UBIQUITIN CONJUGATION,	THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION	UBIQUITIN CONJUGATION,	THIOESTER 2 BOND, LIGASE	UBIQUITIN CONJUGATION UBC7,	UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC7;	UBIQUITIN CONJUGATION, LIGASE, YEAST	UBIQUITIN CONJUGATION UBC2;	UBIQUITIN CONJUGATION,	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN CONJUGATION UBC2;	UBIQUITIN CONJUGATION,	UBIQUITIN-CONJUGATING ENZYME	LIGASE E6AP; UBCH7; BILOBAL	STRUCTURE, ELONGALED SHAPE, E3	CONJUGATING ENZYME	LIGASE E6AP, UBCH7; BILOBAL	STRUCTURE, ELONGATED SHAPE, E3	UBIQUITIN LIGASE, E2 2 UBIQUITIN	CONJUGATING ENCINE	CONTIGATING ENZYME, YEAST	LIGASE UBIQUITIN, UBIQUITIN-	CONJUGATING ENZYME, YEAST	UBIQUITIN-CONJUGATING ENZYME	UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME,
Coumpound		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING	ENZYME; CHAIN: NULL;		UBIQUITIN CONJUGATING	ENZYME; CHAIN: NULL;		UBIQUITIN CONJUGATING	ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATING	ENZYME; CHAIN: NULL;	UBIQUITIN-CONJUGATING	ENZYME RAD6; CHAIN: A, B,	Ċ;	UBIQUITIN-CONJUGATING	ENZYME RAD6; CHAIN: A, B,	ပ်	UBIQUITIN-PROTEIN	LIGASE E3A; CHAIN: A, B, C;	ENZYME E2: CHAIN: D;	UBIQUITIN-PROTEIN	LIGASE E3A; CHAIN: A, B, C;	UBIQUITIN CONJUGATING	ENCYME EZ; CHAIN: D;	UBIQUITIN CONJUGATING	UBIOUITIN CONJUGATING	ENZYME; CHAIN: A;	UBC9, CHAIN: NULL;	
SeqFold score			93.12	81.53								121.4		114.6									104.56				104.4			94.76	
PMF		_				·	_			-							_			_								_			
Verify score		0.38					0.27			0.63							0.73			0.4								0.53			
PSI- BLAST		6.40E-48	6.40E-48	3.20E-44			3.20E-44			3.20E-51		3.20E-51		6.40E-50			6.40E-50			3.20E-42			3.20E-42			3	1.105-53	1.10E-53		4.80E-45	
End		157	148	156			157			157		165		178			178			165			691				991	166		174	
Start		5	9	_			3			9		7		9			9			01			10				/	∞		3	
Chain ID														A			A			D			D				<	V		¥	
PDB ID		2aak	2aak	2e2c			2e2c			2ucz		2ucz		layz			layz			1c4z			1c4z				<u>ş</u>	laca	:	1u9a	
SEQ ID	2	883	883	883			883			883		883		883			883			883			883				883	883		883	

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
										UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
883	1u9a	<	S	173	4.80E-45	8.0	-		UBC9: CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
883	2aak		5	172	8.00E-52	0.62	_		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI, UBIQUITIN CONJUGATION, LIGASE
883	2aak		9	174	8.00E-52			116.78	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
883	2e2c		-	176	1.60E-45			109.63	UBIQUITIN CONJUGATING ENZYME, CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
883	2 e 2c		E.	160	1.60E-45	0.55			UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
883	2ucz		9	160	1.60E-55	0.59			UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7, UBIQUITIN CONJUGATION, LIGASE, YEAST
883	2ucz		1	174	1.60E-55			148.79	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7, UBIQUITIN CONJUGATION, LIGASE, YEAST
883	layz	∢	9	128	4.80E-37	0.19	0.75		UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
883	1c4z	Ω	10	128	1.60E-28	-0.22	0.54		UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP, UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
883	1c4z	Q	10	163	1.60E-28			51.09	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP, UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
883	1qcq	A	8	128	3.20E-39	0.2	9.0		UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
883	1 u9a	V	5	128	3.20E-34	-0.18	0.72		UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME

SEQ ID	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
SO:										UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
883	2nak		5	128	3.20E-38	90.0	0.55		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
883	2aak		9	155	3.20E-38			50.62	UBIQUITIN CONJUGATING ENZYME, CHAIN: NULL,	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
883	2e2c		£	126	1.10E-35	0.17	0.25		UBIQUITIN CONJUGATING ENZYME, CHAIN: NULL,	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
883	2ncz		9	128	1.60E-35	-0.09	0.49		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
883	2ucz		7	136	1.60E-35			56.09	UBIQUITIN CONJUGATING ENZYME, CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
		_								
884	layz	<	9	157	1.10E-45	0.49	-		UBIQUITIN-CONJUGATING ENZYME RAD6: CHAIN: A. B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
884	layz	∢	9	171	1.10E-45			90.32	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
884	lc4z	Ω	01	157	6.40E-39	0.3	_		UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
884	1c4z	۵	01	166	6.40E-39			89.72	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
884	Iqcq	∀	7	168	9.60E-49			87.99	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
884	lqcq	<	∞	157	9.60E-49	9.0	1		UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
884	lu9a	Y	3	168	9.60E-43			74.26	UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME,

PDB Chain Start End PSI- Verify ID AA AA BLAST score	Start End PSI- AA AA BLAST	End PSI- AA BLAST	PSI- BLAST	Ver	ة تِرَ	PMF	SeqFold	Coumpound	PDB annotation
									UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
1u9a A 5 157 9.60E-43 0.17 1	5 157 9.60E-43	157 9.60E-43	9.60E-43	0.17	_			UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
2aak 5 157 6.40E-48 0.38 1	157 6.40E-48	157 6.40E-48	6.40E-48	0.38 1	_	ļ		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
2aak 6 148 6.40E-48 9	148 6.40E-48	148 6.40E-48	6.40E-48	5	5	5	93.12	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
2c2c 1 156 3.20E-44 8	3.20E-44	3.20E-44	3.20E-44			∞ -	81.53	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
2c2c 3 157 3.20E-44 0.27 1	157 3.20E-44	157 3.20E-44	3.20E-44	0.27	-			UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
Zucz 6 157 3.20E-51 0.63 1	157 3.20E-51	157 3.20E-51	3.20E-51	0.63	_			UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
2ucz 7 165 3.20E-51	165 3.20E-51	165 3.20E-51	3.20E-51				121.4	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
layz A 6 178 6.40E-50	6 178 6.40E-50	178 6.40E-50	6.40E-50				114.6	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
layz A 6 178 6.40E-50 0.73 1	6 178 6.40E-50	178 6.40E-50	6.40E-50	 0.73	-			UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
1c4z D 10 165 3.20E-42 0.4 1	10 165 3.20E-42	165 3.20E-42	3.20E-42	0.4		ļ		UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
1c4z D 10 169 3.20E-42	10 169 3.20E-42	169 3.20E-42	3.20E-42				104.56	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP; UBCH7; BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME

10-	PDB	Chain	Start	End	-ISA	Verify	PMF	SeqFold	Coumpound	PDB annotation
=	۵	e	¥¥	*	BLAST	score	score	score		
12	bob	¥	7	166	1.10E-53			104.4	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
1-	1959	A	∞	991	1.10E-53	0.53			UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
ļ —	lu9a	4	3	174	4.80E-45			94.76	UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
1	1u9a	K	8	173	4.80E-45	8.0	-		UBC9; CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
1, ,	2aak		2	172	8.00E-52	0.62	_		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
	2aak		9	174	8.00E-52			116.78	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
	2 e 2c		_	9/1	1.60E-45			109.63	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
	2 e 2c		3	091	1.60E-45	0.55			UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION. UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
1	2ucz		9	160	1.60E-55	0.59	-		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
	Zucz		7	174	1.60E-55			148.79	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7: UBIQUITIN CONJUGATION, LIGASE, YEAST
ł	layz	∢	9	128	4.80E-37	0.19	0.75		UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
	1c4z	Q	01	128	1.60E-28	-0.22	0.54		UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP, UBCH7, BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME

SEQ ID NO:	PDB ID	Chain TD	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
884	1c4z	Ω	01	163	1.60E-28			51.09	UBIQUITIN-PROTEIN LIGASE E3A; CHAIN: A, B, C; UBIQUITIN CONJUGATING ENZYME E2; CHAIN: D;	LIGASE E6AP, UBCH7, BILOBAL STRUCTURE, ELONGATED SHAPE, E3 UBIQUITIN LIGASE, E2 2 UBIQUITIN CONJUGATING ENZYME
884	ldcd	∢	∞	128	3.20E-39	0.2	9.6		UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
884	lu9a	<	\$	128	3.20E-34	-0.18	0.72		UBC9, CHAIN: NULL;	UBIQUITIN-CONJUGATING ENZYME UBIQUITIN-CONJUGATING ENZYME; UBIQUITIN-CONJUGATING ENZYME, UBIQUITIN-DIRECTED 2 PROTEOLYSIS, CELL CYCLE CONTROL, LIGASE
884	2aak		\$	128	3.20E-38	90.0	0.55		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
884	2aak		9	155	3.20E-38			50.62	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC1; UBIQUITIN CONJUGATION, LIGASE
884	2e2c		£	126	1.10E-35	0.17	0.25		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION, UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE
884	2ncz		9	128	1.60E-35	-0.09	0.49		UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
884	2ucz		7	136	1.60E-35			56.09	UBIQUITIN CONTUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBC7; UBIQUITIN CONJUGATION, LIGASE, YEAST
885	lavi	4	237	427	5.40E-07			53.53	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D:	LIPID TRANSPORT APO A-I: LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION
885	lega	٧	39	350	1.60E-37	-0.14	0.01		GTP-BINDING PROTEIN ERA; CHAIN: A. B.	HYDROLASE ERA, GTPASE, RNA- BINDING, RAS-LIKE, HYDROLASE
882	1g7s	4	45	289	6.40E-06	-0.34	0.03		TRANSLATION INITIATION FACTOR IFZ/EIFSB; CHAIN: A:	TRANSLATION TRANSLATIONAL GTPASE
882	lquu	¥.	283	426	1.80E-08	0.23	-0.01		HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A:	CONTRACTILE PROTEIN TRIPLE. HELIX COILED COIL, CONTRACTILE PROTEIN
885	lquu	¥	302	445	1.80E-08	0.23	-0.01		HUMAN SKELETAL MUSCLE	CONTRACTILE PROTEIN TRIPLE-

PDB annotation	THE ANTIGEN COMBINING SITE SUPERANTIGEN FAB VH3 3 SPECIFICITY			IMMUNOGLOBULIN TR 1.9, ANTI- THYROID PEROXIDASE, AUTOANTIBODY, 2 IMMUNOGLOBULIN		DNA-BINDING PROTEIN ISL-1HD DNA-BINDING PROTEIN, HOMEODOMAIN, LIM DOMAIN	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX, TRANSCRIPTION 2 REGULATION	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX, TRANSCRIPTION 2 REGULATION	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX, TRANSCRIPTION 2 REGULATION	COMPLEX (DNA-BINDING PROTEIN/DNA) DNA-BINDING PROTEIN, DNA, PAIRED BOX,
Coumpound	BINDING PROTEIN A; CHAIN: G, H;	IMMUNOGLOBULIN FV FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 1FGV 3 ANTIBODY 'H52' (HUH52-AA FV) 1FGV 4	IMMUNOGLOBULIN IMMUNOGLOBULIN M (IG- M) FV FRAGMENT IIGM 3	TR1.9 FAB; CHAIN: L, H;	IMMUNOGLOBULIN FAB FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 2FGW 3 ANTIBODY 'H52' (HUH52-OZ FAB) 2FGW 4	INSULIN GENE ENHANCER PROTEIN ISL-1; CHAIN: NULL;	PAIRED PROTEIN; CHAIN: A, B. C; DNA; CHAIN: D, E, F	PAIRED PROTEIN; CHAIN: A, B, C; DNA; CHAIN: D, E, F	PAIRED PROTEIN; CHAIN: A, B, C; DNA; CHAIN: D, E, F	PAIRED PROTEIN; CHAIN: A, B, C; DNA; CHAIN: D, E, F
SeqFold score								76.5	72.69	
PMF		0.13	-0.06	0.23	0.1	0.42	_			-
Verify score		0.31	0	0.15	0.32	0.14	0.28			0.02
PSI- BLAST		3.20E-44	1.40E-42	6.40E-41	1.60E-44	9.00E-31	1.10E-28	1.10E-28	9.00E-28	9.00E-28
End		116	125	118	116	275	275	280	273	274
Start		61	61	22	19	216	216	216	217	217
Chain ID			<u>ر</u>	٦	LI .		«	<	В	В
PDB ID		1fgv	ligm	lvge	2fgw	lbw S	<u>1</u> £1	161	161	161
SEQ ID NO:		887	887	887	887	888	888	888	888	888

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PDB annotation	TRANSCRIPTION 2 REGULATION	COMPLEX (GENE REGULATING PROTEIN/DAX, PRD, PAIRED DOMAIN, DNA-BINDING PROTEIN 1PDN 14	COMPLEX (GENE REGULATING PROTEIN/DNA) PAX, PRD, PAIRED DOMAIN, DNA-BINDING PROTEIN IPDN 14	COMPLEX (GENE REGULATING PROTEIN/DNA) PAX, PRD, PAIRED DOMAIN, DNA-BINDING PROTEIN IPDN 14	DNA-BINDING DNA-BINDING, TRANSCRIPTION FACTOR, LFB1/HNF1, 2 HELIX-TURN-HELIX, DNA-BINDING DOMAIN	GENE REGULATION/DNA PAX, PAIRED DOMAIN, TRANSCRIPTION, PROTEIN-DNA INTERACTIONS, 2 GENE REGULATION/DNA	GENE REGULATION/DNA PAX, PAIRED DOMAIN, TRANSCRIPTION, PROTEIN-DNA INTERACTIONS, 2 GENE REGULATION/DNA	GENE REGULATION/DNA PAX, PAIRED DOMAIN, TRANSCRIPTION, PROTEIN-DNA INTERACTIONS, 2 GENE REGULATION/DNA	
Coumpound		PRD PAIRED DOMAIN; CHAIN: C; IPDN 4 DNA; CHAIN: A, B IPDN 5	PRD PAIRED DOMAIN; CHAIN: C; IPDN 4 DNA; CHAIN: A, B IPDN 5	PRD PAIRED DOMAIN; CHAIN: C; IPDN 4 DNA; CHAIN: A, B IPDN 5	LFBI/HNFI TRANSCRIPTION FACTOR; CHAIN: NULL;	HOMEOBOX PROTEIN PAX- 6; CHAIN: A; 26 NUCLEOTIDE DNA; CHAIN: B; 26 NUCLEOTIDE DNA; CHAIN: C;	HOMEOBOX PROTEIN PAX- 6; CHAIN: A; 26 NUCLEOTIDE DNA; CHAIN: B; 26 NUCLEOTIDE DNA; CHAIN: C;	HOMEOBOX PROTEIN PAX- 6; CHAIN: A; 26 NUCLEOTIDE DNA; CHAIN: B; 26 NUCLEOTIDE DNA; CHAIN: C;	
SeqFold score			168.62			143.36			
PMF		 -		-	0.33		_	_	
Verify		0.57		99.0	0.31		0.61	0.48	
PSI- BLAST		9.60E-29	1.10E-63	1.10E-63	1.40E-30	1.60E-67	1.60E-31	1.60E-67	
End AA		152	091	160	280	169	152	169	
Start AA		35	35	35	209	34	35	36	
Chain ID		ပ	U	ပ		∢	∢	⋖	
PDB ID		1pdn	Ipdn	upd1	21fb	брах	xad9	брах	
SEQ EQ		888	888	888	888	888	888	888	

TABLE 6

SEQ ID NO:	Position of Last Amino Acid of Signal Peptide	Maximum Score	Mean Score
445	21	0.993	0.931
446	14	0.975	0.962
447	42	0.986	0.606
448	18	0.908	0.703
449	24	0.967	0.778
450	30	0.992	0.946
452	17	0.997	0.973
454	32	0.907	0.575
455	27	0.931	0.672
456	40	0.988	0.755
457	26	0.986	0.916
458	18	0.920	0.750
459	15	0.946	0.790
460	21	0.993	0.931
461	47	0.942	0.644
463	24	0.886	0.712
464	36	0.985	0.865
465	42	0.965	0.679
466	25	0.980	0.946
467	27	0.969	0.858
469	26	0.950	0.793
470	45	0.983	0.687
471	25	0.981	0.821
472	30	0.998	0.963
473	18	0.977	0.915
474	27	0.949	0.644
475	23	0.913	0.768
476	19	0.947	0.901
477	15	0.936	0.628
478	17	0.956	0.893
479	17	0.942	0.720
480	19	0.952	0.730
481	17	0.970	0.916
483	14	0.975	0.962
486	47	0.955	0.727
488	23	0.991	0.952
495	42	0.986	0.606
496	11	0.971	0.594
502	29	0.896	0.743
509	18	0.908	0.703
510	13	0.959	0.908
512	20	0.957	0.858
516	24	0.967	0.778
517	35	0.991	0.851
518	26	0.939	0.722
519	47	0.983	0.640
522	30	0.992	0.946
538	16	0.974	0.924
550	17	0.997	0.973
551	42	0.947	0.588
555	30	0.981	0.684
576	32	0.907	0.575
577	26	0.973	0.927
578	27	0.931	0.672
589	40	0.988	0.755
590	38	0.985	0.775

505	20	0.938	0.818
595	20	0.920	0.750
611	25	0.949	0.775
616	33	0.995	0.835
617	15	0.946	0.790
623	19	0.921	0.819
627	21	0.993	0.931
634	20	0.961	0.674
635	28	0.954	0.648
645	47	0.942	0.644
647	31	0.962	0.776
650	16	0.949	0.782
651	14	0 963	0.613
654	20	0.984	0.958
670	24	0.886	0.712
673	17	0 934	0.753
678	36	0 985	0.865
695	23	0 954	0.754
707	42	0.965	0.679
708	2	0 979	0.667
709	24	0.984	0.851
710	17	0.911	0.745
717	25	0 980	0.946
718	35	0.988	0.871
726	27	0.969	0.858
730	17	0.981	0.844_
741	22	0.937	0.871
755	17	0.890	0.668
764	26	0.950	0.793
768	32	0.958	0.827
771	45	0.983	0.687
773	39	0.997	0.801
776	17	0.945	0.650
787	32	0.983	0.835
789	25	0.981	0.821 0.815
792	31	0.966	0.813
796	22	0.887	0.691
797	19	0.941	0.963
807	30	0.977	0.903
808	18	0.977	0.915
809	18 27	0.959	0.827
811	16	0.925	0.734
812 815	19	0.934	0.564
816	21	0.960	0.858
818	27	0.949	0.644
821	27	0.943	0.758
823	27	0.908	0.728
833	23	0.913	0.768
837	19	0.947	0.901
841	22	0.967	0.826
845	15	0.936	0.628
846	20	0.975	0.840
851	31	0.985	0.908
852	19	0.965	0.922
853	39	0.984	0.743
857	17	0.956	0.893
858	21	0.957	0.868
861	22	0.975	0.866
868		0.942	0.736
000			1

871	43	0.973	0.560
873	19	0.952	0.730
874	33	0.923	0.717
879	23	0.978	0.911
881	16	0.947	0.884
887	17	0.970	0.916

TABLE 7

SEQ ID NO:	Position of Last Amino Acid of Signal Peptide	Maximum Score	Mean Score
445	21	0.993	0.931
446	14	0.975	0.962
447	42	0.986	0.606
448	18	0.908	0.703
449	24	0.967	0.778
450	30	0.992	0.946
452	17	0.997	0.973
454	32	0.907	0.575
455	27	0.931	0.672
456	40	0.988	0.755
	26	0.986	0.916
457	· · · · · · · · · · · · · · · · · · ·	0.980	0.750
458	18	0.946	0.790
459			0.931
460	21	0.993	0.644
461	47	0.942	
463	24	0.886	0.712
464	36	0.985	0.865
465	42	0.965	0.679
466	25	0.980	0.946
467	27	0.969	0.858
469	26	0.950	0.793
470	45	0.983	0.687
471	25	0.981	0.821
472	30	0.998	0.963
473	18	0.977	0.915
474	27	0.949	0.644
475	23	0.913	0.768
476	19	0.947	0.901
477	15	0.936	0.628
478	17	0.956	0.893
479	17	0.942	0.720
480	19	0.952	0.730
481	17	0.970	0.916
483	14	0.975	0.962
486	47	0.955	0.727
488	23	0.991	0.952
495	42	0.986	0.606
496	11	0.971	0.594
502	29	0.896	0.743
509	18	0.908	0.703
510	13	0.959	0.908
512	20	0.957	0.858
516	24	0.967	0.778
517	35	0.991	0.851
518	26	0.939	0.722
519	47	0.983	0.640
522	30	0.992	0.946
	16	0.992	0.924
538	17	0.974	0.973
550			0.588
551	42	0.947	
555	30	0.981	0.684 0.575
576	32	0.907	
577	26	0.973	0.927
578	27	0.931	0.672
589	40	0.988	0.755

600	38	0.985	0.775
590 595	20	0.938	0.818
611	18	0.920	0.750
615	25	0.949	0.775
616	33	0.995	0.835
	15	0.946	0.790
617	19	0.921	0.819
623	21	0.993	0.931
627	20	0.961	0.674
634	28	0.954	0 648
635	47	0.942	0 644
	31	0.962	0 776
647	16	0.949	0.782
650	14	0.963	0 613
	20	0.984	0.958
654		0.886	0 712
670	24 17	0.934	0 753
673	36	0.985	0.865
678	23	0.954	0.754
695	42	0.965	0.734
707	2	0.979	0.667
708	24	0.979	0.851
	17	0.911	0.745
710	25	0.980	0.946
717	35	0.988	0.871
726	27	0.969	0.858
	17	0.981	0.844
730 741	22	0.937	0.871
755	17	0.890	0.668
764	26	0.950	0.793
768	32	0.958	0.827
771	45	0.983	0.687
773	39	0.997	0.801
776	17	0.945	0.650
787	32	0.983	0.835
789	25	0.981	0.821
792	31	0.966	0.815
796	22	0.887	0.572
797	19	0.941	0.691
807	30	0.998	0.963
808	18	0.977	0.915
809	18	0.977	0.915
811	27	0.959	0.827
812	16	0.925	0.734
815	19	0.934	0.564
816	21	0.960	0.858
818	27	0.949	0.644
821	27	0.943	0.758
823	27	0.908	0.728
833	23	0.913	0.768
837	19	0.947	0.901
841	22	0.967	0.826
845	15	0.936	0.628
846	20	0.975	0.840
851	31	0.985	0.908
852	19	0.965	0.922
853	39	0.984	0.743
857	17	0.956	0.893
858	21	0.957	0.868
861	22	0.975	0.866

868	21	0.942	0.736
871	43	0.973	0.560
873	19	0.952	0.730
874	33	0.923	0.717
879	23	0.978	0.911
881	16	0.947	0.884
887	17	0.970	0.916

TABLE 8

SEQ ID NO: of Nucleotide Sequence	SEQ ID NO: of Polypeptide Sequence	SEQ ID NO: in USSN 09/659,671
1	445	2
2	446	. 5
3	447	6
4	448	7
5	449	8
6	450	9
7	451	11
8	452	12
9	453	13
10	454	14
11	455	15
12	456	16
13	457	17
		18
14	458	
15	459	19
16	460	20
17	461	21
18	462	23
19	463	24
20	464	25
21	465	28
22	466	29
23	467	30
24	468	31
25	469	34
26	470	35
27	471	37
28	472	38
29	473	39
30	474	40
31	475	41
32	476	42
33	477	43
34	478	44
35	479	45
36	480	46
37	481	47
38	482	49
39	483	50
40	484	51
41	485	52
42	486	53
43	487	54
		55
44	488	
45	489	56
46	490	57
47	491	58
48	492	59
49	493	60
50	494	61
51	495	62
52	496	63
53	497	64
54	498	65

55	499	66
56	500	67
57	501	68
58	502	69
59	503	70
60	504	71
61	505	72
62	506	73
63	507	74
64	508	75
65	509	76
66	510	77
67	511	78
68	512	79
69	513	80
70	514	81
71	515	82
72	516	83
73	517	85
74	518	86
75	519	88
76	520	89
77	521	90
78	522	91
79	523	92
80	524	93
81	525	94
82	526	95
83	527	96
84	528	97
85	529	98
86	530	99
87	531	100
88	532	101
89	533	102
90	534	105
91	535	106
92	536	107
93	537	108
94	538	109
95	539	110
96	540	111
97	541 542	112 113
98	542	113
99	544	114
100	545	116
101	546	117
102	547	117
104	548	118
105	549	120
106	550	121
107	551	122
107	552	123
108	553	123
110	554	125
111	555	126
112	556	127
113	557	128
113	558	129
117		

115	559	130
115	560	131
116 117	561	132
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WO 02/22660 PCT/US01/26015

WHAT IS CLAIMED IS:

- 1. An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of SEQ ID NO: 1-444, a mature protein coding portion of SEQ ID NO: 1-444, an active domain coding portion of SEQ ID NO: 1-444, and complementary sequences thereof.
- 2. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide hybridizes to the polynucleotide of claim 1 under stringent hybridization conditions.
- 3. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide has greater than about 90% sequence identity with the polynucleotide of claim 1.
- 4. The polynucleotide of claim 1 wherein said polynucleotide is DNA.
- 5. An isolated polynucleotide of claim 1 wherein said polynucleotide comprises the complementary sequences.
- 6. A vector comprising the polynucleotide of claim 1.
- 7. An expression vector comprising the polynucleotide of claim 1.
- 8. A host cell genetically engineered to comprise the polynucleotide of claim 1.
- 9. A host cell genetically engineered to comprise the polynucleotide of claim 1 operatively associated with a regulatory sequence that modulates expression of the polynucleotide in the host cell.
- 10. An isolated polypeptide, wherein the polypeptide is selected from the group consisting of:
 - (a) a polypeptide encoded by any one of the polynucleotides of claim 1; and
 - (b) a polypeptide encoded by a polynucleotide hybridizing under stringent conditions with any one of SEQ ID NO: 1-444.

- 11. A composition comprising the polypeptide of claim 10 and a carrier.
- 12. An antibody directed against the polypeptide of claim 10.
- 13. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polynucleotide of claim 1 for a period sufficient to form the complex; and
- b) detecting the complex, so that if a complex is detected, the polynucleotide of claim 1 is detected.
- 14. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample under stringent hybridization conditions with nucleic acid primers that anneal to the polynucleotide of claim 1 under such conditions;
- b) amplifying a product comprising at least a portion of the polynucleotide of claim 1; and
- c) detecting said product and thereby the polynucleotide of claim 1 in the sample.
- 15. The method of claim 14, wherein the polynucleotide is an RNA molecule and the method further comprises reverse transcribing an annealed RNA molecule into a cDNA polynucleotide.
- 16. A method for detecting the polypeptide of claim 10 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex; and
- b) detecting formation of the complex, so that if a complex formation is detected, the polypeptide of claim 10 is detected.
- 17. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10 under conditions sufficient to form a polypeptide/compound complex; and
- b) detecting the complex, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.

- 18. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10, in a cell, under conditions sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and
- b) detecting the complex by detecting reporter gene sequence expression, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.
- 19. A method of producing the polypeptide of claim 10, comprising,
- a) culturing a host cell comprising a polynucleotide sequence selected from SEQ ID NO: 1-444, a mature protein coding portion of SEQ ID NO: 1-444, an active domain coding portion of SEQ ID NO: 1-444, complementary sequences thereof and a polynucleotide sequence hybridizing under stringent conditions to SEQ ID NO: 1-444, under conditions sufficient to express the polypeptide in said cell; and
 - b) isolating the polypeptide from the cell culture or cells of step (a).
- 20. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of any one of the polypeptides SEQ ID NO: 217-432, or 649-864, the mature protein portion thereof, or the active domain thereof.
- 21. The polypeptide of claim 20 wherein the polypeptide is provided on a polypeptide array.
- 22. A collection of polynucleotides, wherein the collection comprising the sequence information of at least one of SEQ ID NO: 1-444.
- 23. The collection of claim 22, wherein the collection is provided on a nucleic acid array.
- 24. The collection of claim 23, wherein the array detects full-matches to any one of the polynucleotides in the collection.
- 25. The collection of claim 23, wherein the array detects mismatches to any one of the polynucleotides in the collection.

- 26. The collection of claim 22, wherein the collection is provided in a computer-readable format.
- 27. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.
- 28. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising an antibody that specifically binds to a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.

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(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization International Bureau



(43) International Publication Date 21 March 2002 (21.03.2002)

PCT

(10) International Publication Number WO 02/22660 A2

(51) International Patent Classification?: C07K 14/00

(21) International Application Number: PCT/US01/26015

(22) International Filing Date:

10 September 2001 (10.09.2001)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

09/659,671

11 September 2000 (11.09.2000) U

(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application:

US

09/659,671 (CIP)

Filed on

11 September 2000 (11.09.2000)

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- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GII, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- without international search report and to be republished upon receipt of that report
- with sequence listing part of description published separately in electronic form and available upon request from the International Bureau

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

/22660 A

(54) Title: NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

(57) Abstract: The present invention provides nucleic acids, novel polypeptide sequences encoded by these nucleic acids and uses thereof

The compositions of the present invention additionally include vectors, including expression vectors, containing the polynucleotides of the invention, cells genetically engineered to contain such polynucleotides and cells genetically engineered to express such polynucleotides.

The present invention relates to a collection or library of at least one novel nucleic acid sequence assembled from expressed sequence tags (ESTs) isolated mainly by sequencing by hybridization (SBH), and in some cases, sequences obtained from one or more public databases. The invention relates also to the proteins encoded by such polynucleotides, along with therapeutic, diagnostic and research utilities for these polynucleotides and proteins. These nucleic acid sequences are designated as SEQ ID NO: 1-444. The polypeptides sequences are designated SEQ ID NO: 445-888. The nucleic acids and polypeptides are provided in the Sequence Listing. In the nucleic acids provided in the Sequence Listing, A is adenosine; C is cytosine; G is guanine; T is thymine; and N is unknown or any of the four bases.

The nucleic acid sequences of the present invention also include, nucleic acid sequences that hybridize to the complement of SEQ ID NO: 1-444 under stringent hybridization conditions; nucleic acid sequences which are allelic variants or species homologues of any of the nucleic acid sequences recited above, or nucleic acid sequences that encode a peptide comprising a specific domain or truncation of the peptides encoded by SEQ ID NO: 1-444. A polynucleotide comprising a nucleotide sequence having at least 90% identity to an identifying sequence of SEQ ID NO: 1-444 or a degenerate variant or fragment thereof. The identifying sequence can be 100 base pairs in length.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO: 1-444. The sequence information can be a segment of any one of SEQ ID NO: 1-444 that uniquely identifies or represents the sequence information of SEQ ID NO: 1-444.

A collection as used in this application can be a collection of only one polynucleotide. The collection of sequence information or identifying information of each sequence can be provided on a nucleic acid array. In one embodiment, segments of sequence information are provided on a nucleic acid array to detect the polynucleotide that contains the segment. The array can be designed to detect full-match or mismatch to the polynucleotide that contains the segment. The collection can also be provided in a computer-readable format.

This invention also includes the reverse or direct complement of any of the nucleic acid sequences recited above; cloning or expression vectors containing the nucleic acid sequences; and host cells or organisms transformed with these expression vectors. Nucleic acid sequences (or their reverse or direct complements) according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology, such as use as hybridization

probes, use as primers for PCR, use in an array, use in computer-readable media, use in sequencing full-length genes, use for chromosome and gene mapping, use in the recombinant production of protein, and use in the generation of anti-sense DNA or RNA, their chemical analogs and the like.

In a preferred embodiment, the nucleic acid sequences of SEQ ID NO: 1-444 or novel segments or parts of the nucleic acids of the invention are used as primers in expression assays that are well known in the art. In a particularly preferred embodiment, the nucleic acid sequences of SEQ ID NO: 1-444 or novel segments or parts of the nucleic acids provided herein are used in diagnostics for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The isolated polynucleotides of the invention include, but are not limited to, a polynucleotide comprising any one of the nucleotide sequences set forth in SEQ ID NO: 1-444; a polynucleotide comprising any of the full length protein coding sequences of SEQ ID NO: 1-444; and a polynucleotide comprising any of the nucleotide sequences of the mature protein coding sequences of SEQ ID NO: 1-444. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent hybridization conditions to (a) the complement of any one of the nucleotide sequences set forth in SEQ ID NO: 1-444; (b) a nucleotide sequence encoding any one of the amino acid sequences set forth in the Sequence Listing; (c) a polynucleotide which is an allelic variant of any polynucleotides recited above; (d) a polynucleotide which encodes a species homolog (e.g. orthologs) of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of any of the polypeptides comprising an amino acid sequence set forth in the Sequence Listing.

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising any of the amino acid sequences set forth in SEQ ID NO: 445-888; or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides with biological activity that are encoded by (a) any of the polynucleotides having a nucleotide sequence set forth in SEQ ID NO: 1-444; or (b) polynucleotides that hybridize to the complement of the polynucleotides of (a) under stringent hybridization conditions. Biologically or immunologically active variants of any of the polypeptide sequences in the Sequence Listing, and "substantial equivalents" thereof (e.g., with at least about 65%, 70%, 75%, 80%, 85%, 90%, 95%, 98% or 99% amino acid sequence identity) that preferably retain biological activity are also contemplated. The polypeptides of the invention may be wholly or partially chemically synthesized but are preferably produced by recombinant means using the genetically engineered cells (e.g. host cells) of the invention.

The invention also provides compositions comprising a polypeptide of the invention. Polypeptide compositions of the invention may further comprise an acceptable carrier, such as a hydrophilic, e.g., pharmaceutically acceptable, carrier.

The invention also provides host cells transformed or transfected with a polynucleotide of the invention.

The invention also relates to methods for producing a polypeptide of the invention comprising growing a culture of the host cells of the invention in a suitable culture medium under conditions permitting expression of the desired polypeptide, and purifying the polypeptide from the culture or from the host cells. Preferred embodiments include those in which the protein produced by such process is a mature form of the protein.

Polynucleotides according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology. These techniques include use as hybridization probes, use as oligomers, or primers, for PCR, use for chromosome and gene mapping, use in the recombinant production of protein, and use in generation of anti-sense DNA or RNA, their chemical analogs and the like. For example, when the expression of an mRNA is largely restricted to a particular cell or tissue type, polynucleotides of the invention can be used as hybridization probes to detect the presence of the particular cell or tissue mRNA in a sample using, e.g., in situ hybridization.

In other exemplary embodiments, the polynucleotides are used in diagnostics as expressed sequence tags for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The polypeptides according to the invention can be used in a variety of conventional procedures and methods that are currently applied to other proteins. For example, a polypeptide of the invention can be used to generate an antibody that specifically binds the polypeptide. Such antibodies, particularly monoclonal antibodies, are useful for detecting or quantitating the polypeptide in tissue. The polypeptides of the invention can also be used as molecular weight markers, and as a food supplement.

Methods are also provided for preventing, treating, or ameliorating a medical condition which comprises the step of administering to a mammalian subject a therapeutically effective amount of a composition comprising a polypeptide of the present invention and a pharmaceutically acceptable carrier.

In particular, the polypeptides and polynucleotides of the invention can be utilized, for example, in methods for the prevention and/or treatment of disorders involving aberrant protein expression or biological activity.

The present invention further relates to methods for detecting the presence of the polynucleotides or polypeptides of the invention in a sample. Such methods can, for example, be utilized as part of prognostic and diagnostic evaluation of disorders as recited herein and for the identification of subjects exhibiting a predisposition to such conditions. The invention provides a method for detecting the polynucleotides of the invention in a sample, comprising contacting the sample with a compound that binds to and forms a complex with the polynucleotide of interest for a period sufficient to form the complex and under conditions sufficient to form a complex and detecting the complex such that if a complex is detected, the polynucleotide of interest is detected. The invention also provides a method for detecting the polypeptides of the invention in a sample comprising contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex and detecting the formation of the complex such that if a complex is formed, the polypeptide is detected.

The invention also provides kits comprising polynucleotide probes and/or monoclonal antibodies, and optionally quantitative standards, for carrying out methods of the invention. Furthermore, the invention provides methods for evaluating the efficacy of drugs, and monitoring the progress of patients, involved in clinical trials for the treatment of disorders as recited above.

The invention also provides methods for the identification of compounds that modulate (i.e., increase or decrease) the expression or activity of the polynucleotides and/or polypeptides of the invention. Such methods can be utilized, for example, for the identification of compounds that can ameliorate symptoms of disorders as recited herein. Such methods can include, but are not limited to, assays for identifying compounds and other substances that interact with (e.g., bind to) the polypeptides of the invention. The invention provides a method for identifying a compound that binds to the polypeptides of the invention comprising contacting the compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and detecting the complex by detecting the reporter gene sequence expression such that if expression of the reporter gene is detected the compound the binds to a polypeptide of the invention is identified.

The methods of the invention also provide methods for treatment which involve the administration of the polynucleotides or polypeptides of the invention to individuals exhibiting symptoms or tendencies. In addition, the invention encompasses methods for treating diseases or disorders as recited herein comprising administering compounds and other substances that modulate the overall activity of the target gene products. Compounds and other substances can

effect such modulation either on the level of target gene/protein expression or target protein activity.

The polypeptides of the present invention and the polynucleotides encoding them are also useful for the same functions known to one of skill in the art as the polypeptides and polynucleotides to which they have homology (set forth in Table 2); for which they have a signature region (as set forth in Table 3); or for which they have homology to a gene family (as set forth in Table 4). If no homology is set forth for a sequence, then the polypeptides and polynucleotides of the present invention are useful for a variety of applications, as described herein, including use in arrays for detection.

4. DETAILED DESCRIPTION OF THE INVENTION

4.1 DEFINITIONS

It must be noted that as used herein and in the appended claims, the singular forms "a", "an" and "the" include plural references unless the context clearly dictates otherwise.

The term "active" refers to those forms of the polypeptide which retain the biologic and/or immunologic activities of any naturally occurring polypeptide. According to the invention, the terms "biologically active" or "biological activity" refer to a protein or peptide having structural, regulatory or biochemical functions of a naturally occurring molecule. Likewise "immunologically active" or "immunological activity" refers to the capability of the natural, recombinant or synthetic polypeptide to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies.

The term "activated cells" as used in this application are those cells which are engaged in extracellular or intracellular membrane trafficking, including the export of secretory or enzymatic molecules as part of a normal or disease process.

The terms "complementary" or "complementarity" refer to the natural binding of polynucleotides by base pairing. For example, the sequence 5'-AGT-3' binds to the complementary sequence 3'-TCA-5'. Complementarity between two single-stranded molecules may be "partial" such that only some of the nucleic acids bind or it may be "complete" such that total complementarity exists between the single stranded molecules. The degree of complementarity between the nucleic acid strands has significant effects on the efficiency and strength of the hybridization between the nucleic acid strands.

The term "embryonic stem cells (ES)" refers to a cell that can give rise to many differentiated cell types in an embryo or an adult, including the germ cells. The term "germ line stem cells (GSCs)" refers to stem cells derived from primordial stem cells that provide a steady

and continuous source of germ cells for the production of gametes. The term "primordial germ cells (PGCs)" refers to a small population of cells set aside from other cell lineages particularly from the yolk sac, mesenteries, or gonadal ridges during embryogenesis that have the potential to differentiate into germ cells and other cells. PGCs are the source from which GSCs and ES cells are derived. The PGCs, the GSCs and the ES cells are capable of self-renewal. Thus these cells not only populate the germ line and give rise to a plurality of terminally differentiated cells that comprise the adult specialized organs, but are able to regenerate themselves.

The term "expression modulating fragment," EMF, means a series of nucleotides which modulates the expression of an operably linked ORF or another EMF.

As used herein, a sequence is said to "modulate the expression of an operably linked sequence" when the expression of the sequence is altered by the presence of the EMF. EMFs include, but are not limited to, promoters, and promoter modulating sequences (inducible elements). One class of EMFs are nucleic acid fragments which induce the expression of an operably linked ORF in response to a specific regulatory factor or physiological event.

The terms "nucleotide sequence" or "nucleic acid" or "polynucleotide" or "oligonculeotide" are used interchangeably and refer to a heteropolymer of nucleotides or the sequence of these nucleotides. These phrases also refer to DNA or RNA of genomic or synthetic origin which may be single-stranded or double-stranded and may represent the sense or the antisense strand, to peptide nucleic acid (PNA) or to any DNA-like or RNA-like material. In the sequences herein A is adenine, C is cytosine, T is thymine, G is guanine and N is A, C, G or T (U). It is contemplated that where the polynucleotide is RNA, the T (thymine) in the sequences provided herein is substituted with U (uracil). Generally, nucleic acid segments provided by this invention may be assembled from fragments of the genome and short oligonucleotide linkers, or from a series of oligonucleotides, or from individual nucleotides, to provide a synthetic nucleic acid which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial or viral operon, or a eukaryotic gene.

The terms "oligonucleotide fragment" or a "polynucleotide fragment", "portion," or "segment" or "probe" or "primer" are used interchangeably and refer to a sequence of nucleotide residues which are at least about 5 nucleotides, more preferably at least about 7 nucleotides, more preferably at least about 11 nucleotides and most preferably at least about 17 nucleotides. The fragment is preferably less than about 500 nucleotides, preferably less than about 200 nucleotides, more preferably less than about 100 nucleotides, more preferably less than about 50 nucleotides and most preferably less than 30 nucleotides. Preferably the probe is from about 6 nucleotides to about 200 nucleotides, preferably from about 15 to about 50 nucleotides, more preferably from about 17 to 30

nucleotides and most preferably from about 20 to 25 nucleotides. Preferably the fragments can be used in polymerase chain reaction (PCR), various hybridization procedures or microarray procedures to identify or amplify identical or related parts of mRNA or DNA molecules. A fragment or segment may uniquely identify each polynucleotide sequence of the present invention. Preferably the fragment comprises a sequence substantially similar to any one of SEQ ID NO: 1-444.

Probes may, for example, be used to determine whether specific mRNA molecules are present in a cell or tissue or to isolate similar nucleic acid sequences from chromosomal DNA as described by Walsh et al. (Walsh, P.S. et al., 1992, PCR Methods Appl 1:241-250). They may be labeled by nick translation, Klenow fill-in reaction, PCR, or other methods well known in the art. Probes of the present invention, their preparation and/or labeling are elaborated in Sambrook, J. et al., 1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY; or Ausubel, F.M. et al., 1989, Current Protocols in Molecular Biology, John Wiley & Sons, New York NY, both of which are incorporated herein by reference in their entirety.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO: 1-444. The sequence information can be a segment of any one of SEQ ID NO: 1-444 that uniquely identifies or represents the sequence information of that sequence of SEQ ID NO: 1-444. One such segment can be a twenty-mer nucleic acid sequence because the probability that a twenty-mer is fully matched in the human genome is 1 in 300. In the human genome, there are three billion base pairs in one set of chromosomes. Because 4²⁰ possible twenty-mers exist, there are 300 times more twenty-mers than there are base pairs in a set of human chromosomes. Using the same analysis, the probability for a seventeen-mer to be fully matched in the human genome is approximately 1 in 5. When these segments are used in arrays for expression studies, fifteen-mer segments can be used. The probability that the fifteen-mer is fully matched in the expressed sequences is also approximately one in five because expressed sequences comprise less than approximately 5% of the entire genome sequence.

Similarly, when using sequence information for detecting a single mismatch, a segment can be a twenty-five mer. The probability that the twenty-five mer would appear in a human genome with a single mismatch is calculated by multiplying the probability for a full match $(1 \div 4^{25})$ times the increased probability for mismatch at each nucleotide position (3×25) . The probability that an eighteen mer with a single mismatch can be detected in an array for expression studies is approximately one in five. The probability that a twenty-mer with a single mismatch can be detected in a human genome is approximately one in five.

The term "open reading frame," ORF, means a series of nucleotide triplets coding for amino acids without any termination codons and is a sequence translatable into protein.

The terms "operably linked" or "operably associated" refer to functionally related nucleic acid sequences. For example, a promoter is operably associated or operably linked with a coding sequence if the promoter controls the transcription of the coding sequence. While operably linked nucleic acid sequences can be contiguous and in the same reading frame, certain genetic elements e.g. repressor genes are not contiguously linked to the coding sequence but still control transcription/translation of the coding sequence.

The term "pluripotent" refers to the capability of a cell to differentiate into a number of differentiated cell types that are present in an adult organism. A pluripotent cell is restricted in its differentiation capability in comparison to a totipotent cell.

The terms "polypeptide" or "peptide" or "amino acid sequence" refer to an oligopeptide, peptide, polypeptide or protein sequence or fragment thereof and to naturally occurring or synthetic molecules. A polypeptide "fragment," "portion," or "segment" is a stretch of amino acid residues of at least about 5 amino acids, preferably at least about 7 amino acids, more preferably at least about 9 amino acids and most preferably at least about 17 or more amino acids. The peptide preferably is not greater than about 500 amino acids, more preferably less than 200 amino acids more preferably less than 150 amino acids and most preferably less than 100 amino acids. Preferably the peptide is from about 5 to about 200 amino acids. To be active, any polypeptide must have sufficient length to display biological and/or immunological activity.

The term "naturally occurring polypeptide" refers to polypeptides produced by cells that have not been genetically engineered and specifically contemplates various polypeptides arising from post-translational modifications of the polypeptide including, but not limited to, acetylation, carboxylation, glycosylation, phosphorylation, lipidation and acylation.

The term "translated protein coding portion" means a sequence which encodes for the full length protein which may include any leader sequence or any processing sequence.

The term "mature protein coding sequence" means a sequence which encodes a peptide or protein without a signal or leader sequence. The "mature protein portion" means that portion of the protein which does not include a signal or leader sequence. The peptide may have been produced by processing in the cell which removes any leader/signal sequence. The mature protein portion may or may not include an initial methionine residue. The methionine residue may be removed from the protein during processing in the cell. The peptide may be produced synthetically or the protein may have been produced using a polynucleotide only encoding for the mature protein coding sequence.

The term "derivative" refers to polypeptides chemically modified by such techniques as ubiquitination, labeling (e.g., with radionuclides or various enzymes), covalent polymer attachment such as pegylation (derivatization with polyethylene glycol) and insertion or substitution by chemical synthesis of amino acids such as ornithine, which do not normally occur in human proteins.

The term "variant" (or "analog") refers to any polypeptide differing from naturally occurring polypeptides by amino acid insertions, deletions, and substitutions, created using, e.g., recombinant DNA techniques. Guidance in determining which amino acid residues may be replaced, added or deleted without abolishing activities of interest, may be found by comparing the sequence of the particular polypeptide with that of homologous peptides and minimizing the number of amino acid sequence changes made in regions of high homology (conserved regions) or by replacing amino acids with consensus sequence.

Alternatively, recombinant variants encoding these same or similar polypeptides may be synthesized or selected by making use of the "redundancy" in the genetic code. Various codon substitutions, such as the silent changes which produce various restriction sites, may be introduced to optimize cloning into a plasmid or viral vector or expression in a particular prokaryotic or eukaryotic system. Mutations in the polypucleotide sequence may be reflected in the polypeptide or domains of other peptides added to the polypeptide to modify the properties of any part of the polypeptide, to change characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate.

Preferably, amino acid "substitutions" are the result of replacing one amino acid with another amino acid having similar structural and/or chemical properties, i.e., conservative amino acid replacements. "Conservative" amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, and/or the amphipathic nature of the residues involved. For example, nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan, and methionine; polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine, and glutamine; positively charged (basic) amino acids include arginine, lysine, and histidine; and negatively charged (acidic) amino acids include aspartic acid and glutamic acid. "Insertions" or "deletions" are preferably in the range of about 1 to 20 amino acids, more preferably 1 to 10 amino acids. The variation allowed may be experimentally determined by systematically making insertions, deletions, or substitutions of amino acids in a polypeptide molecule using recombinant DNA techniques and assaying the resulting recombinant variants for activity.

Alternatively, where alteration of function is desired, insertions, deletions or non-conservative alterations can be engineered to produce altered polypeptides. Such alterations

can, for example, alter one or more of the biological functions or biochemical characteristics of the polypeptides of the invention. For example, such alterations may change polypeptide characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate. Further, such alterations can be selected so as to generate polypeptides that are better suited for expression, scale up and the like in the host cells chosen for expression. For example, cysteine residues can be deleted or substituted with another amino acid residue in order to eliminate disulfide bridges.

The terms "purified" or "substantially purified" as used herein denotes that the indicated nucleic acid or polypeptide is present in the substantial absence of other biological macromolecules, e.g., polynucleotides, proteins, and the like. In one embodiment, the polynucleotide or polypeptide is purified such that it constitutes at least 95% by weight, more preferably at least 99% by weight, of the indicated biological macromolecules present (but water, buffers, and other small molecules, especially molecules having a molecular weight of less than 1000 daltons, can be present).

The term "isolated" as used herein refers to a nucleic acid or polypeptide separated from at least one other component (e.g., nucleic acid or polypeptide) present with the nucleic acid or polypeptide in its natural source. In one embodiment, the nucleic acid or polypeptide is found in the presence of (if anything) only a solvent, buffer, ion, or other component normally present in a solution of the same. The terms "isolated" and "purified" do not encompass nucleic acids or polypeptides present in their natural source.

The term "recombinant," when used herein to refer to a polypeptide or protein, means that a polypeptide or protein is derived from recombinant (e.g., microbial, insect, or mammalian) expression systems. "Microbial" refers to recombinant polypeptides or proteins made in bacterial or fungal (e.g., yeast) expression systems. As a product, "recombinant microbial" defines a polypeptide or protein essentially free of native endogenous substances and unaccompanied by associated native glycosylation. Polypeptides or proteins expressed in most bacterial cultures, e.g., E. coli, will be free of glycosylation modifications; polypeptides or proteins expressed in yeast will have a glycosylation pattern in general different from those expressed in mammalian cells.

The term "recombinant expression vehicle or vector" refers to a plasmid or phage or virus or vector, for expressing a polypeptide from a DNA (RNA) sequence. An expression vehicle can comprise a transcriptional unit comprising an assembly of (1) a genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers, (2) a structural or coding sequence which is transcribed into mRNA and translated into protein, and (3) appropriate transcription initiation and termination sequences. Structural units intended for use

in yeast or eukaryotic expression systems preferably include a leader sequence enabling extracellular secretion of translated protein by a host cell. Alternatively, where recombinant protein is expressed without a leader or transport sequence, it may include an amino terminal methionine residue. This residue may or may not be subsequently cleaved from the expressed recombinant protein to provide a final product.

The term "recombinant expression system" means host cells which have stably integrated a recombinant transcriptional unit into chromosomal DNA or carry the recombinant transcriptional unit extrachromosomally. Recombinant expression systems as defined herein will express heterologous polypeptides or proteins upon induction of the regulatory elements linked to the DNA segment or synthetic gene to be expressed. This term also means host cells which have stably integrated a recombinant genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers. Recombinant expression systems as defined herein will express polypeptides or proteins endogenous to the cell upon induction of the regulatory elements linked to the endogenous DNA segment or gene to be expressed. The cells can be prokaryotic or eukaryotic.

The term "secreted" includes a protein that is transported across or through a membrane, including transport as a result of signal sequences in its amino acid sequence when it is expressed in a suitable host cell. "Secreted" proteins include without limitation proteins secreted wholly (e.g., soluble proteins) or partially (e.g., receptors) from the cell in which they are expressed. "Secreted" proteins also include without limitation proteins that are transported across the membrane of the endoplasmic reticulum. "Secreted" proteins are also intended to include proteins containing non-typical signal sequences (e.g. Interleukin-1 Beta, see Krasney, P.A. and Young, P.R. (1992) Cytokine 4(2): 134 -143) and factors released from damaged cells (e.g. Interleukin-1 Receptor Antagonist, see Arend, W.P. et. al. (1998) Annu. Rev. Immunol. 16:27-55)

Where desired, an expression vector may be designed to contain a "signal or leader sequence" which will direct the polypeptide through the membrane of a cell. Such a sequence may be naturally present on the polypeptides of the present invention or provided from heterologous protein sources by recombinant DNA techniques.

The term "stringent" is used to refer to conditions that are commonly understood in the art as stringent. Stringent conditions can include highly stringent conditions (i.e., hybridization to filter-bound DNA in 0.5 M NaHPO₄, 7% sodium dodecyl sulfate (SDS), 1 mM EDTA at 65°C, and washing in 0.1X SSC/0.1% SDS at 68°C), and moderately stringent conditions (i.e., washing in 0.2X SSC/0.1% SDS at 42°C). Other exemplary hybridization conditions are described herein in the examples.

In instances of hybridization of deoxyoligonucleotides, additional exemplary stringent hybridization conditions include washing in 6X SSC/0.05% sodium pyrophosphate at 37°C (for 14-base oligonucleotides), 48°C (for 17-base oligos), 55°C (for 20-base oligonucleotides), and 60°C (for 23-base oligonucleotides).

As used herein, "substantially equivalent" can refer both to nucleotide and amino acid sequences, for example a mutant sequence, that varies from a reference sequence by one or more substitutions, deletions, or additions, the net effect of which does not result in an adverse functional dissimilarity between the reference and subject sequences. Typically, such a substantially equivalent sequence varies from one of those listed herein by no more than about 35% (i.e., the number of individual residue substitutions, additions, and/or deletions in a substantially equivalent sequence, as compared to the corresponding reference sequence, divided by the total number of residues in the substantially equivalent sequence is about 0.35 or less). Such a sequence is said to have 65% sequence identity to the listed sequence. In one embodiment, a substantially equivalent, e.g., mutant, sequence of the invention varies from a listed sequence by no more than 30% (70% sequence identity); in a variation of this embodiment, by no more than 25% (75% sequence identity); and in a further variation of this embodiment, by no more than 20% (80% sequence identity) and in a further variation of this embodiment, by no more than 10% (90% sequence identity) and in a further variation of this embodiment, by no more that 5% (95% sequence identity). Substantially equivalent, e.g., mutant, amino acid sequences according to the invention preferably have at least 80% sequence identity with a listed amino acid sequence, more preferably at least 85% sequence identity, more preferably at least 90% sequence identity, more preferably at least 95% identity, more preferably at least 98% identity, and most preferably at least 99% identity. Substantially equivalent nucleotide sequences of the invention can have lower percent sequence identities, taking into account, for example, the redundancy or degeneracy of the genetic code. Preferably, nucleotide sequence has at least about 65% identity, more preferably at least about 75% identity, more preferably at least about 80% sequence identity, more preferably at least about 85% sequence identity, more preferably at least about 90% sequence identity, and most preferably at least about 95% identity, more preferably at least about 98% sequence identity, and most preferably at least about 99% sequence identity. For the purposes of the present invention, sequences having substantially equivalent biological activity and substantially equivalent expression characteristics are considered substantially equivalent. For the purposes of determining equivalence, truncation of the mature sequence (e.g., via a mutation which creates a spurious stop codon) should be disregarded. Sequence identity may be determined, e.g., using the Jotun Hein method (Hein, J.

(1990) Methods Enzymol. 183:626-645). Identity between sequences can also be determined by other methods known in the art, e.g. by varying hybridization conditions.

The term "totipotent" refers to the capability of a cell to differentiate into all of the cell types of an adult organism.

The term "transformation" means introducing DNA into a suitable host cell so that the DNA is replicable, either as an extrachromosomal element, or by chromosomal integration. The term "transfection" refers to the taking up of an expression vector by a suitable host cell, whether or not any coding sequences are in fact expressed. The term "infection" refers to the introduction of nucleic acids into a suitable host cell by use of a virus or viral vector.

As used herein, an "uptake modulating fragment," UMF, means a series of nucleotides which mediate the uptake of a linked DNA fragment into a cell. UMFs can be readily identified using known UMFs as a target sequence or target motif with the computer-based systems described below. The presence and activity of a UMF can be confirmed by attaching the suspected UMF to a marker sequence. The resulting nucleic acid molecule is then incubated with an appropriate host under appropriate conditions and the uptake of the marker sequence is determined. As described above, a UMF will increase the frequency of uptake of a linked marker sequence.

Each of the above terms is meant to encompass all that is described for each, unless the context dictates otherwise.

4.2 NUCLEIC ACIDS OF THE INVENTION

Nucleotide sequences of the invention are set forth in the Sequence Listing.

The isolated polynucleotides of the invention include a polynucleotide comprising the nucleotide sequences of SEQ ID NO: 1-444; a polynucleotide encoding any one of the peptide sequences of SEQ ID NO: 445-888; and a polynucleotide comprising the nucleotide sequence encoding the mature protein coding sequence of the polypeptides of any one of SEQ ID NO: 445-888. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent conditions to (a) the complement of any of the nucleotides sequences of SEQ ID NO: 1-444; (b) nucleotide sequences encoding any one of the amino acid sequences set forth in the Sequence Listing as SEQ ID NO: 445-888; (c) a polynucleotide which is an allelic variant of any polynucleotide recited above; (d) a polynucleotide which encodes a species homolog of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of the polypeptides of SEQ ID NO: 445-888. Domains of interest may depend on the nature of the encoded polypeptide; e.g., domains in receptor-like polypeptides include ligand-binding,

extracellular, transmembrane, or cytoplasmic domains, or combinations thereof; domains in immunoglobulin-like proteins include the variable immunoglobulin-like domains; domains in enzyme-like polypeptides include catalytic and substrate binding domains; and domains in ligand polypeptides include receptor-binding domains.

The polynucleotides of the invention include naturally occurring or wholly or partially synthetic DNA, e.g., cDNA and genomic DNA, and RNA, e.g., mRNA. The polynucleotides may include all of the coding region of the cDNA or may represent a portion of the coding region of the cDNA.

The present invention also provides genes corresponding to the cDNA sequences disclosed herein. The corresponding genes can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include the preparation of probes or primers from the disclosed sequence information for identification and/or amplification of genes in appropriate genomic libraries or other sources of genomic materials. Further 5' and 3' sequence can be obtained using methods known in the art. For example, full length cDNA or genomic DNA that corresponds to any of the polynucleotides of SEQ ID NO: 1-444 can be obtained by screening appropriate cDNA or genomic DNA libraries under suitable hybridization conditions using any of the polynucleotides of SEQ ID NO: 1-444 or a portion thereof as a probe. Alternatively, the polynucleotides of SEQ ID NO: 1-444 may be used as the basis for suitable primer(s) that allow identification and/or amplification of genes in appropriate genomic DNA or cDNA libraries.

The nucleic acid sequences of the invention can be assembled from ESTs and sequences (including cDNA and genomic sequences) obtained from one or more public databases, such as dbEST, gbpri, and UniGene. The EST sequences can provide identifying sequence information, representative fragment or segment information, or novel segment information for the full-length gene.

The polynucleotides of the invention also provide polynucleotides including nucleotide sequences that are substantially equivalent to the polynucleotides recited above. Polynucleotides according to the invention can have, e.g., at least about 65%, at least about 70%, at least about 75%, at least about 80%, 81%, 82%, 83%, 84%, more typically at least about 85%, 86%, 87%, 88%, 89%, more typically at least about 90%, 91%, 92%, 93%, 94%, and even more typically at least about 95%, 96%, 97%, 98%, 99%, sequence identity to a polynucleotide recited above.

Included within the scope of the nucleic acid sequences of the invention are nucleic acid sequence fragments that hybridize under stringent conditions to any of the nucleotide sequences of SEQ ID NO: 1-444, or complements thereof, which fragment is greater than about 5 nucleotides, preferably 7 nucleotides, more preferably greater than 9 nucleotides and most preferably greater than 17 nucleotides. Fragments of, e.g. 15, 17, or 20 nucleotides or more that

are selective for (i.e. specifically hybridize to) any one of the polynucleotides of the invention are contemplated. Probes capable of specifically hybridizing to a polynucleotide can differentiate polynucleotide sequences of the invention from other polynucleotide sequences in the same family of genes or can differentiate human genes from genes of other species, and are preferably based on unique nucleotide sequences.

The sequences falling within the scope of the present invention are not limited to these specific sequences, but also include allelic and species variations thereof. Allelic and species variations can be routinely determined by comparing the sequence provided in SEQ ID NO: 1-444, a representative fragment thereof, or a nucleotide sequence at least 90% identical, preferably 95% identical, to SEQ ID NO: 1-444 with a sequence from another isolate of the same species. Furthermore, to accommodate codon variability, the invention includes nucleic acid molecules coding for the same amino acid sequences as do the specific ORFs disclosed herein. In other words, in the coding region of an ORF, substitution of one codon for another codon that encodes the same amino acid is expressly contemplated.

The nearest neighbor or homology result for the nucleic acids of the present invention, including SEQ ID NO: 1-444, can be obtained by searching a database using an algorithm or a program. Preferably, a BLAST which stands for Basic Local Alignment Search Tool is used to search for local sequence alignments (Altshul, S.F. J Mol. Evol. 36 290-300 (1993) and Altschul S.F. et al. J. Mol. Biol. 21:403-410 (1990)). Alternatively a FASTA version 3 search against Genpept, using Fastxy algorithm.

Species homologs (or orthologs) of the disclosed polynucleotides and proteins are also provided by the present invention. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source from the desired species.

The invention also encompasses allelic variants of the disclosed polynucleotides or proteins; that is, naturally-occurring alternative forms of the isolated polynucleotide which also encode proteins which are identical, homologous or related to that encoded by the polynucleotides.

The nucleic acid sequences of the invention are further directed to sequences which encode variants of the described nucleic acids. These amino acid sequence variants may be prepared by methods known in the art by introducing appropriate nucleotide changes into a native or variant polynucleotide. There are two variables in the construction of amino acid sequence variants: the location of the mutation and the nature of the mutation. Nucleic acids encoding the amino acid sequence variants are preferably constructed by mutating the polynucleotide to encode an amino acid sequence that does not occur in nature. These nucleic

acid alterations can be made at sites that differ in the nucleic acids from different species (variable positions) or in highly conserved regions (constant regions). Sites at such locations will typically be modified in series, e.g., by substituting first with conservative choices (e.g., hydrophobic amino acid to a different hydrophobic amino acid) and then with more distant choices (e.g., hydrophobic amino acid to a charged amino acid), and then deletions or insertions may be made at the target site. Amino acid sequence deletions generally range from about 1 to 30 residues, preferably about 1 to 10 residues, and are typically contiguous. Amino acid insertions include amino- and/or carboxyl-terminal fusions ranging in length from one to one hundred or more residues, as well as intrasequence insertions of single or multiple amino acid residues. Intrasequence insertions may range generally from about 1 to 10 amino residues, preferably from 1 to 5 residues. Examples of terminal insertions include the heterologous signal sequences necessary for secretion or for intracellular targeting in different host cells and sequences such as FLAG or poly-histidine sequences useful for purifying the expressed protein.

In a preferred method, polynucleotides encoding the novel amino acid sequences are changed via site-directed mutagenesis. This method uses oligonucleotide sequences to alter a polynucleotide to encode the desired amino acid variant, as well as sufficient adjacent nucleotides on both sides of the changed amino acid to form a stable duplex on either side of the site of being changed. In general, the techniques of site-directed mutagenesis are well known to those of skill in the art and this technique is exemplified by publications such as, Edelman et al., DNA 2:183 (1983). A versatile and efficient method for producing site-specific changes in a polynucleotide sequence was published by Zoller and Smith, Nucleic Acids Res. 10:6487-6500 (1982). PCR may also be used to create amino acid sequence variants of the novel nucleic acids. When small amounts of template DNA are used as starting material, primer(s) that differs slightly in sequence from the corresponding region in the template DNA can generate the desired amino acid variant. PCR amplification results in a population of product DNA fragments that differ from the polynucleotide template encoding the polypeptide at the position specified by the primer. The product DNA fragments replace the corresponding region in the plasmid and this gives a polynucleotide encoding the desired amino acid variant.

A further technique for generating amino acid variants is the cassette mutagenesis technique described in Wells et al., *Gene* 34:315 (1985); and other mutagenesis techniques well known in the art, such as, for example, the techniques in Sambrook et al., supra, and *Current Protocols in Molecular Biology*, Ausubel et al. Due to the inherent degeneracy of the genetic code, other DNA sequences which encode substantially the same or a functionally equivalent amino acid sequence may be used in the practice of the invention for the cloning and expression

of these novel nucleic acids. Such DNA sequences include those which are capable of hybridizing to the appropriate novel nucleic acid sequence under stringent conditions.

Polynucleotides encoding preferred polypeptide truncations of the invention can be used to generate polynucleotides encoding chimeric or fusion proteins comprising one or more domains of the invention and heterologous protein sequences.

The polynucleotides of the invention additionally include the complement of any of the polynucleotides recited above. The polynucleotide can be DNA (genomic, cDNA, amplified, or synthetic) or RNA. Methods and algorithms for obtaining such polynucleotides are well known to those of skill in the art and can include, for example, methods for determining hybridization conditions that can routinely isolate polynucleotides of the desired sequence identities.

In accordance with the invention, polynucleotide sequences comprising the mature protein coding sequences corresponding to any one of SEQ ID NO: 1-444, or functional equivalents thereof, may be used to generate recombinant DNA molecules that direct the expression of that nucleic acid, or a functional equivalent thereof, in appropriate host cells. Also included are the cDNA inserts of any of the clones identified herein.

A polynucleotide according to the invention can be joined to any of a variety of other nucleotide sequences by well-established recombinant DNA techniques (see Sambrook J et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY). Useful nucleotide sequences for joining to polynucleotides include an assortment of vectors, e.g., plasmids, cosmids, lambda phage derivatives, phagemids, and the like, that are well known in the art. Accordingly, the invention also provides a vector including a polynucleotide of the invention and a host cell containing the polynucleotide. In general, the vector contains an origin of replication functional in at least one organism, convenient restriction endonuclease sites, and a selectable marker for the host cell. Vectors according to the invention include expression vectors, replication vectors, probe generation vectors, and sequencing vectors. A host cell according to the invention can be a prokaryotic or eukaryotic cell and can be a unicellular organism or part of a multicellular organism.

The present invention further provides recombinant constructs comprising a nucleic acid having any of the nucleotide sequences of SEQ ID NO: 1-444 or a fragment thereof or any other polynucleotides of the invention. In one embodiment, the recombinant constructs of the present invention comprise a vector, such as a plasmid or viral vector, into which a nucleic acid having any of the nucleotide sequences of SEQ ID NO: 1-444 or a fragment thereof is inserted, in a forward or reverse orientation. In the case of a vector comprising one of the ORFs of the present invention, the vector may further comprise regulatory sequences, including for example, a promoter, operably linked to the ORF. Large numbers of suitable vectors and promoters are

known to those of skill in the art and are commercially available for generating the recombinant constructs of the present invention. The following vectors are provided by way of example. Bacterial: pBs, phagescript, PsiX174, pBluescript SK, pBs KS, pNH8a, pNH16a, pNH18a, pNH46a (Stratagene); pTrc99A, pKK223-3, pKK233-3, pDR540, pRIT5 (Pharmacia). Eukaryotic: pWLneo, pSV2cat, pOG44, PXTI, pSG (Stratagene) pSVK3, pBPV, pMSG, pSVL (Pharmacia).

The isolated polynucleotide of the invention may be operably linked to an expression control sequence such as the pMT2 or pED expression vectors disclosed in Kaufman et al., Nucleic Acids Res. 19, 4485-4490 (1991), in order to produce the protein recombinantly. Many suitable expression control sequences are known in the art. General methods of expressing recombinant proteins are also known and are exemplified in R. Kaufman, Methods in Enzymology 185, 537-566 (1990). As defined herein "operably linked" means that the isolated polynucleotide of the invention and an expression control sequence are situated within a vector or cell in such a way that the protein is expressed by a host cell which has been transformed (transfected) with the ligated polynucleotide/expression control sequence.

Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, and trc. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein-I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art. Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, e.g., the ampicillin resistance gene of E. coli and S. cerevisiae TRP1 gene, and a promoter derived from a highly-expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3-phosphoglycerate kinase (PGK), a-factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated protein into the periplasmic space or extracellular medium. Optionally, the heterologous sequence can encode a fusion protein including an amino terminal identification peptide imparting desired characteristics, e.g., stabilization or simplified purification of expressed recombinant product. Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or

more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and to, if desirable, provide amplification within the host. Suitable prokaryotic hosts for transformation include *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium* and various species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*, although others may also be employed as a matter of choice.

As a representative but non-limiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and GEM 1 (Promega Biotech, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed. Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter is induced or derepressed by appropriate means (e.g., temperature shift or chemical induction) and cells are cultured for an additional period. Cells are typically harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract retained for further purification.

Polynucleotides of the invention can also be used to induce immune responses. For example, as described in Fan et al., *Nat. Biotech.* 17:870-872 (1999), incorporated herein by reference, nucleic acid sequences encoding a polypeptide may be used to generate antibodies against the encoded polypeptide following topical administration of naked plasmid DNA or following injection, and preferably intramuscular injection of the DNA. The nucleic acid sequences are preferably inserted in a recombinant expression vector and may be in the form of naked DNA.

4.3 ANTISENSE NUCLEIC ACIDS

Another aspect of the invention pertains to isolated antisense nucleic acid molecules that are hybridizable to or complementary to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NO: 1-444, or fragments, analogs or derivatives thereof. An "antisense" nucleic acid comprises a nucleotide sequence that is complementary to a "sense" nucleic acid encoding a protein, *e.g.*, complementary to the coding strand of a double-stranded cDNA molecule or complementary to an mRNA sequence. In specific aspects, antisense nucleic acid molecules are provided that comprise a sequence complementary to at least about 10, 25, 50, 100, 250 or 500 nucleotides or an entire coding strand, or to only a portion thereof. Nucleic acid molecules encoding fragments, homologs, derivatives and analogs of a protein of any of SEQ ID

PCT/US01/26015 WO 02/22660

NO: 445-888 or antisense nucleic acids complementary to a nucleic acid sequence of SEQ ID NO: 1-444 are additionally provided.

In one embodiment, an antisense nucleic acid molecule is antisense to a "coding region" of the coding strand of a nucleotide sequence of the invention. The term "coding region" refers to the region of the nucleotide sequence comprising codons which are translated into amino acid residues. In another embodiment, the antisense nucleic acid molecule is antisense to a "noncoding region" of the coding strand of a nucleotide sequence of the invention. The term "noncoding region" refers to 5' and 3' sequences which flank the coding region that are not translated into amino acids (i.e., also referred to as 5' and 3' untranslated regions).

Given the coding strand sequences encoding a nucleic acid disclosed herein (e.g., SEQ ID NO: 1-444), antisense nucleic acids of the invention can be designed according to the rules of Watson and Crick or Hoogsteen base pairing. The antisense nucleic acid molecule can be complementary to the entire coding region of an mRNA, but more preferably is an oligonucleotide that is antisense to only a portion of the coding or noncoding region of a mRNA. For example, the antisense oligonucleotide can be complementary to the region surrounding the translation start site of a mRNA. An antisense oligonucleotide can be, for example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis or enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (e.g., an antisense oligonucleotide) can be chemically synthesized using naturally occurring nucleotides or variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids, e.g., phosphorothioate derivatives and acridine substituted nucleotides can be used.

Examples of modified nucleotides that can be used to generate the antisense nucleic acid include: 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxylmethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the

antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (*i.e.*, RNA transcribed from the inserted nucleic acid will be of an antisense orientation to a target nucleic acid of interest, described further in the following subsection).

The antisense nucleic acid molecules of the invention are typically administered to a subject or generated in situ such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding a protein according to the invention to thereby inhibit expression of the protein, e.g., by inhibiting transcription and/or translation. The hybridization can be by conventional nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule that binds to DNA duplexes, through specific interactions in the major groove of the double helix. An example of a route of administration of antisense nucleic acid molecules of the invention includes direct injection at a tissue site. Alternatively, antisense nucleic acid molecules can be modified to target selected cells and then administered systemically. For example, for systemic administration, antisense molecules can be modified such that they specifically bind to receptors or antigens expressed on a selected cell surface, e.g., by linking the antisense nucleic acid molecules to peptides or antibodies that bind to cell surface receptors or antigens. The antisense nucleic acid molecules can also be delivered to cells using the vectors described herein. To achieve sufficient intracellular concentrations of antisense molecules, vector constructs in which the antisense nucleic acid molecule is placed under the control of a strong pol II or pol III promoter are preferred.

In yet another embodiment, the antisense nucleic acid molecule of the invention is an α -anomeric nucleic acid molecule. An α -anomeric nucleic acid molecule forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual β -units, the strands run parallel to each other (Gaultier *et al.* (1987) *Nucleic Acids Res* 15: 6625-6641). The antisense nucleic acid molecule can also comprise a 2'-o-methylribonucleotide (Inoue *et al.* (1987) *Nucleic Acids Res* 15: 6131-6148) or a chimeric RNA -DNA analogue (Inoue *et al.* (1987) *FEBS Lett* 215: 327-330).

4.4 RIBOZYMES AND PNA MOIETIES

In still another embodiment, an antisense nucleic acid of the invention is a ribozyme. Ribozymes are catalytic RNA molecules with ribonuclease activity that are capable of cleaving a single-stranded nucleic acid, such as an mRNA, to which they have a complementary region. Thus, ribozymes (e.g., hammerhead ribozymes (described in Haselhoff and Gerlach (1988) *Nature* 334:585-591)) can be used to catalytically cleave a mRNA transcripts to thereby inhibit translation of a mRNA. A ribozyme having specificity for a nucleic acid of the invention can be

designed based upon the nucleotide sequence of a DNA disclosed herein (i.e., SEQ ID NO: 1-444). For example, a derivative of a Tetrahymena L-19 IVS RNA can be constructed in which the nucleotide sequence of the active site is complementary to the nucleotide sequence to be cleaved in an mRNA of SEQ ID NO: 1-444 (see, e.g., Cech et al. U.S. Pat. No. 4,987,071; and Cech et al. U.S. Pat. No. 5,116,742). Alternatively, polynucleotides of the invention can be used to select a catalytic RNA having a specific ribonuclease activity from a pool of RNA molecules. See, e.g., Bartel et al., (1993) Science 261:1411-1418.

Alternatively, gene expression can be inhibited by targeting nucleotide sequences complementary to the regulatory region (e.g., promoter and/or enhancers) to form triple helical structures that prevent transcription of the gene in target cells. See generally, Helene. (1991) Anticancer Drug Des. 6: 569-84; Helene. et al. (1992) Ann. N.Y. Acad. Sci. 660:27-36; and Maher (1992) Bioassays 14: 807-15.

In various embodiments, the nucleic acids of the invention can be modified at the base moiety, sugar moiety or phosphate backbone to improve, e.g., the stability, hybridization, or solubility of the molecule. For example, the deoxyribose phosphate backbone of the nucleic acids can be modified to generate peptide nucleic acids (see Hyrup et al. (1996) Bioorg Med Chem 4: 5-23). As used herein, the terms "peptide nucleic acids" or "PNAs" refer to nucleic acid mimics, e.g., DNA mimics, in which the deoxyribose phosphate backbone is replaced by a pseudopeptide backbone and only the four natural nucleobases are retained. The neutral backbone of PNAs has been shown to allow for specific hybridization to DNA and RNA under conditions of low ionic strength. The synthesis of PNA oligomers can be performed using standard solid phase peptide synthesis protocols as described in Hyrup et al. (1996) above; Perry-O'Keefe et al. (1996) PNAS 93: 14670-675.

PNAs of the invention can be used in therapeutic and diagnostic applications. For example, PNAs can be used as antisense or antigene agents for sequence-specific modulation of gene expression by, e.g., inducing transcription or translation arrest or inhibiting replication. PNAs of the invention can also be used, e.g., in the analysis of single base pair mutations in a gene by, e.g., PNA directed PCR clamping; as artificial restriction enzymes when used in combination with other enzymes, e.g., S1 nucleases (Hyrup B. (1996) above); or as probes or primers for DNA sequence and hybridization (Hyrup et al. (1996), above; Perry-O'Keefe (1996), above).

In another embodiment, PNAs of the invention can be modified, e.g., to enhance their stability or cellular uptake, by attaching lipophilic or other helper groups to PNA, by the formation of PNA-DNA chimeras, or by the use of liposomes or other techniques of drug delivery known in the art. For example, PNA-DNA chimeras can be generated that may

combine the advantageous properties of PNA and DNA. Such chimeras allow DNA recognition enzymes, e.g., RNase H and DNA polymerases, to interact with the DNA portion while the PNA portion would provide high binding affinity and specificity. PNA-DNA chimeras can be linked using linkers of appropriate lengths selected in terms of base stacking, number of bonds between the nucleobases, and orientation (Hyrup (1996) above). The synthesis of PNA-DNA chimeras can be performed as described in Hyrup (1996) above and Finn et al. (1996) Nucl Acids Res 24: 3357-63. For example, a DNA chain can be synthesized on a solid support using standard phosphoramidite coupling chemistry, and modified nucleoside analogs, e.g., 5'-(4-methoxytrityl)amino-5'-deoxy-thymidine phosphoramidite, can be used between the PNA and the 5' end of DNA (Mag et al. (1989) Nucl Acid Res 17: 5973-88). PNA monomers are then coupled in a stepwise manner to produce a chimeric molecule with a 5' PNA segment and a 3' DNA segment (Finn et al. (1996) above). Alternatively, chimeric molecules can be synthesized with a 5' DNA segment and a 3' PNA segment. See, Petersen et al. (1975) Bioorg Med Chem Lett 5: 1119-11124.

In other embodiments, the oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors in vivo), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84:648-652; PCT Publication No. W088/09810) or the blood-brain barrier (see, e.g., PCT Publication No. W089/10134). In addition, oligonucleotides can be modified with hybridization triggered cleavage agents (See, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents. (See, e.g., Zon, 1988, Pharm. Res. 5: 539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, a hybridization triggered cross-linking agent, a transport agent, a hybridization-triggered cleavage agent, etc.

4.5 HOSTS

The present invention further provides host cells genetically engineered to contain the polynucleotides of the invention. For example, such host cells may contain nucleic acids of the invention introduced into the host cell using known transformation, transfection or infection methods. The present invention still further provides host cells genetically engineered to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell.

Knowledge of nucleic acid sequences allows for modification of cells to permit, or increase, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous

recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous promoter so that the cells express the polypeptide at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the encoding sequences. See, for example, PCT International Publication No. WO94/12650, PCT International Publication No. WO92/20808, and PCT International Publication No. WO91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the coding sequence, amplification of the marker DNA by standard selection methods results in coamplification of the desired protein coding sequences in the cells.

The host cell can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Introduction of the recombinant construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, or electroporation (Davis, L. et al., Basic Methods in Molecular Biology (1986)). The host cells containing one of the polynucleotides of the invention, can be used in conventional manners to produce the gene product encoded by the isolated fragment (in the case of an ORF) or can be used to produce a heterologous protein under the control of the EMF.

Any host/vector system can be used to express one or more of the ORFs of the present invention. These include, but are not limited to, eukaryotic hosts such as HeLa cells, Cv-1 cell, COS cells, 293 cells, and Sf9 cells, as well as prokaryotic host such as *E. coli* and *B. suhtilis*. The most preferred cells are those which do not normally express the particular polypeptide or protein or which expresses the polypeptide or protein at low natural level. Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described by Sambrook, et al., in Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, New York (1989), the disclosure of which is hereby incorporated by reference.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts, described by Gluzman, Cell 23:175 (1981). Other cell lines capable of expressing a compatible vector are, for example, the C127, monkey COS cells, Chinese Hamster Ovary (CHO) cells, human kidney 293 cells, human epidermal A431 cells, human Colo205 cells, 3T3

cells, CV-1 cells, other transformed primate cell lines, normal diploid cells, cell strains derived from *in vitro* culture of primary tissue, primary explants, HeLa cells, mouse L cells, BHK, HL-60, U937, HaK or Jurkat cells. Mammalian expression vectors will comprise an origin of replication, a suitable promoter and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking nontranscribed sequences. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early promoter, enhancer, splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements. Recombinant polypeptides and proteins produced in bacterial culture are usually isolated by initial extraction from cell pellets, followed by one or more salting-out, aqueous ion exchange or size exclusion chromatography steps. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps. Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents.

Alternatively, it may be possible to produce the protein in lower eukaryotes such as yeast or insects or in prokaryotes such as bacteria. Potentially suitable yeast strains include Saccharomyces cerevisiae, Schizosaccharomyces pombe, Kluyveromyces strains, Candida, or any yeast strain capable of expressing heterologous proteins. Potentially suitable bacterial strains include Escherichia coli, Bacillus subtilis, Salmonella typhimurium, or any bacterial strain capable of expressing heterologous proteins. If the protein is made in yeast or bacteria, it may be necessary to modify the protein produced therein, for example by phosphorylation or glycosylation of the appropriate sites, in order to obtain the functional protein. Such covalent attachments may be accomplished using known chemical or enzymatic methods.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequence include polyadenylation signals, mRNA stability elements, splice

sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the host cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

4.6 POLYPEPTIDES OF THE INVENTION

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising: the amino acid sequences set forth as any one of SEQ ID NO: 445-888 or an amino acid sequence encoded by any one of the nucleotide sequences SEQ ID NO: 1-444 or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides preferably with biological or immunological activity that are encoded by: (a) a polynucleotide having any one of the nucleotide sequences set forth in SEQ ID NO: 1-444 or (b)

polynucleotides encoding any one of the amino acid sequences set forth as SEQ ID NO: 445-888 or (c) polynucleotides that hybridize to the complement of the polynucleotides of either (a) or (b) under stringent hybridization conditions. The invention also provides biologically active or immunologically active variants of any of the amino acid sequences set forth as SEQ ID NO: 445-888 or the corresponding full length or mature protein; and "substantial equivalents" thereof (e.g., with at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, 86%, 87%, 88%, 89%, at least about 90%, 91%, 92%, 93%, 94%, typically at least about 95%, 96%, 97%, more typically at least about 98%, or most typically at least about 99% amino acid identity) that retain biological activity. Polypeptides encoded by allelic variants may have a similar, increased, or decreased activity compared to polypeptides comprising SEQ ID NO: 445-888.

Fragments of the proteins of the present invention which are capable of exhibiting biological activity are also encompassed by the present invention. Fragments of the protein may be in linear form or they may be cyclized using known methods, for example, as described in H. U. Saragovi, et al., Bio/Technology 10, 773-778 (1992) and in R. S. McDowell, et al., J. Amer. Chem. Soc. 114, 9245-9253 (1992), both of which are incorporated herein by reference. Such fragments may be fused to carrier molecules such as immunoglobulins for many purposes, including increasing the valency of protein binding sites.

The present invention also provides both full-length and mature forms (for example, without a signal sequence or precursor sequence) of the disclosed proteins. The protein coding sequence is identified in the sequence listing by translation of the disclosed nucleotide sequences. The mature form of such protein may be obtained by expression of a full-length polynucleotide in a suitable mammalian cell or other host cell. The sequence of the mature form of the protein is also determinable from the amino acid sequence of the full-length form. Where proteins of the present invention are membrane bound, soluble forms of the proteins are also provided. In such forms, part or all of the regions causing the proteins to be membrane bound are deleted so that the proteins are fully secreted from the cell in which they are expressed.

Protein compositions of the present invention may further comprise an acceptable carrier, such as a hydrophilic, e.g., pharmaceutically acceptable, carrier.

The present invention further provides isolated polypeptides encoded by the nucleic acid fragments of the present invention or by degenerate variants of the nucleic acid fragments of the present invention. By "degenerate variant" is intended nucleotide fragments which differ from a nucleic acid fragment of the present invention (e.g., an ORF) by nucleotide sequence but, due to the degeneracy of the genetic code, encode an identical polypeptide sequence. Preferred nucleic acid fragments of the present invention are the ORFs that encode proteins.

A variety of methodologies known in the art can be utilized to obtain any one of the isolated polypeptides or proteins of the present invention. At the simplest level, the amino acid sequence can be synthesized using commercially available peptide synthesizers. The synthetically-constructed protein sequences, by virtue of sharing primary, secondary or tertiary structural and/or conformational characteristics with proteins may possess biological properties in common therewith, including protein activity. This technique is particularly useful in producing small peptides and fragments of larger polypeptides. Fragments are useful, for example, in generating antibodies against the native polypeptide. Thus, they may be employed as biologically active or immunological substitutes for natural, purified proteins in screening of therapeutic compounds and in immunological processes for the development of antibodies.

The polypeptides and proteins of the present invention can alternatively be purified from cells which have been altered to express the desired polypeptide or protein. As used herein, a cell is said to be altered to express a desired polypeptide or protein when the cell, through genetic manipulation, is made to produce a polypeptide or protein which it normally does not produce or which the cell normally produces at a lower level. One skilled in the art can readily adapt procedures for introducing and expressing either recombinant or synthetic sequences into eukaryotic or prokaryotic cells in order to generate a cell which produces one of the polypeptides or proteins of the present invention.

The invention also relates to methods for producing a polypeptide comprising growing a culture of host cells of the invention in a suitable culture medium, and purifying the protein from the cells or the culture in which the cells are grown. For example, the methods of the invention include a process for producing a polypeptide in which a host cell containing a suitable expression vector that includes a polynucleotide of the invention is cultured under conditions that allow expression of the encoded polypeptide. The polypeptide can be recovered from the culture, conveniently from the culture medium, or from a lysate prepared from the host cells and further purified. Preferred embodiments include those in which the protein produced by such process is a full length or mature form of the protein.

In an alternative method, the polypeptide or protein is purified from bacterial cells which naturally produce the polypeptide or protein. One skilled in the art can readily follow known methods for isolating polypeptides and proteins in order to obtain one of the isolated polypeptides or proteins of the present invention. These include, but are not limited to, immunochromatography, HPLC, size-exclusion chromatography, ion-exchange chromatography, and immuno-affinity chromatography. See, e.g., Scopes, Protein Purification: Principles and Practice, Springer-Verlag (1994); Sambrook, et al., in Molecular Cloning: A Laboratory Manual; Ausubel et al., Current Protocols in Molecular Biology. Polypeptide fragments that

retain biological/immunological activity include fragments comprising greater than about 100 amino acids, or greater than about 200 amino acids, and fragments that encode specific protein domains.

The purified polypeptides can be used in *in vitro* binding assays which are well known in the art to identify molecules which bind to the polypeptides. These molecules include but are not limited to, for e.g., small molecules, molecules from combinatorial libraries, antibodies or other proteins. The molecules identified in the binding assay are then tested for antagonist or agonist activity in *in vivo* tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

In addition, the peptides of the invention or molecules capable of binding to the peptides may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for SEQ ID NO: 445-888.

The protein of the invention may also be expressed as a product of transgenic animals, e.g., as a component of the milk of transgenic cows, goats, pigs, or sheep which are characterized by somatic or germ cells containing a nucleotide sequence encoding the protein.

The proteins provided herein also include proteins characterized by amino acid sequences similar to those of purified proteins but into which modification are naturally provided or deliberately engineered. For example, modifications, in the peptide or DNA sequence, can be made by those skilled in the art using known techniques. Modifications of interest in the protein sequences may include the alteration, substitution, replacement, insertion or deletion of a selected amino acid residue in the coding sequence. For example, one or more of the cysteine residues may be deleted or replaced with another amino acid to alter the conformation of the molecule. Techniques for such alteration, substitution, replacement, insertion or deletion are well known to those skilled in the art (see, e.g., U.S. Pat. No. 4,518,584). Preferably, such alteration, substitution, replacement, insertion or deletion retains the desired activity of the protein. Regions of the protein that are important for the protein function can be determined by various methods known in the art including the alanine-scanning method which involved systematic substitution of single or strings of amino acids with alanine, followed by testing the resulting alanine-containing variant for biological activity. This type of analysis determines the importance of the substituted amino acid(s) in biological activity. Regions of the protein that are important for protein function may be determined by the eMATRIX program.

Other fragments and derivatives of the sequences of proteins which would be expected to retain protein activity in whole or in part and are useful for screening or other immunological

methodologies may also be easily made by those skilled in the art given the disclosures herein... Such modifications are encompassed by the present invention.

The protein may also be produced by operably linking the isolated polynucleotide of the invention to suitable control sequences in one or more insect expression vectors, and employing an insect expression system. Materials and methods for baculovirus/insect cell expression systems are commercially available in kit form from, e.g., Invitrogen, San Diego, Calif., U.S.A. (the MaxBatTM kit), and such methods are well known in the art, as described in Summers and Smith, Texas Agricultural Experiment Station Bulletin No. 1555 (1987), incorporated herein by reference. As used herein, an insect cell capable of expressing a polynucleotide of the present invention is "transformed."

The protein of the invention may be prepared by culturing transformed host cells under culture conditions suitable to express the recombinant protein. The resulting expressed protein may then be purified from such culture (i.e., from culture medium or cell extracts) using known purification processes, such as gel filtration and ion exchange chromatography. The purification of the protein may also include an affinity column containing agents which will bind to the protein; one or more column steps over such affinity resins as concanavalin A-agarose, heparin-toyopearlTM or Cibacrom blue 3GA SepharoseTM; one or more steps involving hydrophobic interaction chromatography using such resins as phenyl ether, butyl ether, or propyl ether; or immunoaffinity chromatography.

Alternatively, the protein of the invention may also be expressed in a form which will facilitate purification. For example, it may be expressed as a fusion protein, such as those of maltose binding protein (MBP), glutathione-S-transferase (GST) or thioredoxin (TRX), or as a His tag. Kits for expression and purification of such fusion proteins are commercially available from New England BioLab (Beverly, Mass.), Pharmacia (Piscataway, N.J.) and Invitrogen, respectively. The protein can also be tagged with an epitope and subsequently purified by using a specific antibody directed to such epitope. One such epitope ("FLAG®") is commercially available from Kodak (New Haven, Conn.).

Finally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, e.g., silica gel having pendant methyl or other aliphatic groups, can be employed to further purify the protein. Some or all of the foregoing purification steps, in various combinations, can also be employed to provide a substantially homogeneous isolated recombinant protein. The protein thus purified is substantially free of other mammalian proteins and is defined in accordance with the present invention as an "isolated protein."

The polypeptides of the invention include analogs (variants). This embraces fragments, as well as peptides in which one or more amino acids has been deleted, inserted, or substituted. Also, analogs of the polypeptides of the invention embrace fusions of the polypeptides or modifications of the polypeptides of the invention, wherein the polypeptide or analog is fused to another moiety or moieties, e.g., targeting moiety or another therapeutic agent. Such analogs may exhibit improved properties such as activity and/or stability. Examples of moieties which may be fused to the polypeptide or an analog include, for example, targeting moieties which provide for the delivery of polypeptide to pancreatic cells, e.g., antibodies to pancreatic cells, antibodies to immune cells such as T-cells, monocytes, dendritic cells, granulocytes, etc., as well as receptor and ligands expressed on pancreatic or immune cells. Other moieties which may be fused to the polypeptide include therapeutic agents which are used for treatment, for example, immunosuppressive drugs such as cyclosporin, SK506, azathioprine, CD3 antibodies and steroids. Also, polypeptides may be fused to immune modulators, and other cytokines such as alpha or beta interferon.

4.6.1 DETERMINING POLYPEPTIDE AND POLYNUCLEOTIDE IDENTITY AND SIMILARITY

Preferred identity and/or similarity are designed to give the largest match between the sequences tested. Methods to determine identity and similarity are codified in computer programs including, but are not limited to, the GCG program package, including GAP (Devereux, J., et al., Nucleic Acids Research 12(1):387 (1984); Genetics Computer Group, University of Wisconsin, Madison, WI), BLASTP, BLASTN, BLASTX, FASTA (Altschul, S.F. et al., J. Molec. Biol. 215:403-410 (1990), PSI-BLAST (Altschul S.F. et al., Nucleic Acids Res. vol. 25, pp. 3389-3402, herein incorporated by reference), eMatrix software (Wu et al., J. Comp. Biol., Vol. 6, pp. 219-235 (1999), herein incorporated by reference), eMotif software (Nevill-Manning et al, ISMB-97, Vol. 4, pp. 202-209, herein incorporated by reference), pFam software (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1), pp. 320-322 (1998), herein incorporated by reference), the GeneAtlas software (Molecular Simulations Inc. (MSI), San Diego, CA) (Sanchez and Sali (1998) Proc. Natl. Acad. Sci., 95, 13597-13602; Kitson DH et al, (2000) "Remote homology detection using structural modeling - an evaluation" Submitted; Fischer and Eisenberg (1996) Protein Sci. 5, 947-955), Neural Network SignalP V1.1 program (from Center for Biological Sequence Analysis, The Technical University of Denmark), and the Kyte-Doolittle hydrophobocity prediction algorithm (J. Mol Biol, 157, pp. 105-31 (1982), incorporated herein by reference). The BLAST programs are publicly available from the National Center for Biotechnology Information (NCBI) and other sources (BLAST Manual,

Altschul, S., et al. NCB NLM NIH Bethesda, MD 20894; Altschul, S., et al., J. Mol. Biol. 215:403-410 (1990).

4.7 CHIMERIC AND FUSION PROTEINS

The invention also provides chimeric or fusion proteins. As used herein, a "chimeric protein" or "fusion protein" comprises a polypeptide of the invention operatively linked to another polypeptide. Within a fusion protein the polypeptide according to the invention can correspond to all or a portion of a protein according to the invention. In one embodiment, a fusion protein comprises at least one biologically active portion of a protein according to the invention. In another embodiment, a fusion protein comprises at least two biologically active portions of a protein according to the invention. Within the fusion protein, the term "operatively linked" is intended to indicate that the polypeptide according to the invention and the other polypeptide are fused in-frame to each other. The polypeptide can be fused to the N-terminus or C-terminus.

For example, in one embodiment a fusion protein comprises a polypeptide according to the invention operably linked to the extracellular domain of a second protein.

In another embodiment, the fusion protein is a GST-fusion protein in which the polypeptide sequences of the invention are fused to the C-terminus of the GST (i.e., glutathione S-transferase) sequences.

In another embodiment, the fusion protein is an immunoglobulin fusion protein in which the polypeptide sequences according to the invention comprise one or more domains fused to sequences derived from a member of the immunoglobulin protein family. The immunoglobulin fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject to inhibit an interaction between a ligand and a protein of the invention on the surface of a cell, to thereby suppress signal transduction *in vivo*. The immunoglobulin fusion proteins can be used to affect the bioavailability of a cognate ligand. Inhibition of the ligand/protein interaction may be useful therapeutically for both the treatment of proliferative and differentiative disorders, *e.g.*, cancer as well as modulating (*e.g.*, promoting or inhibiting) cell survival. Moreover, the immunoglobulin fusion proteins of the invention can be used as immunogens to produce antibodies in a subject, to purify ligands, and in screening assays to identify molecules that inhibit the interaction of a polypeptide of the invention with a ligand.

A chimeric or fusion protein of the invention can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, e.g., by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive ends as appropriate, alkaline phosphatase treatment to

avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers that give rise to complementary overhangs between two consecutive gene fragments that can subsequently be annealed and reamplified to generate a chimeric gene sequence (see, for example, Ausubel et al. (eds.) Current Protocols in Molecular Biology, John Wiley & Sons, 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST polypeptide). A nucleic acid encoding a polypeptide of the invention can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the protein of the invention.

4.8 GENE THERAPY

Mutations in the polynucleotides of the invention may result in loss of normal function of the encoded protein. The invention thus provides gene therapy to restore normal activity of the polypeptides of the invention; or to treat disease states involving polypeptides of the invention. Delivery of a functional gene encoding polypeptides of the invention to appropriate cells is effected ex vivo, in situ, or in vivo by use of vectors, and more particularly viral vectors (e.g., adenovirus, adeno-associated virus, or a retrovirus), or ex vivo by use of physical DNA transfer methods (e.g., liposomes or chemical treatments). See, for example, Anderson, Nature, supplement to vol. 392, no. 6679, pp.25-20 (1998). For additional reviews of gene therapy technology see Friedmann, Science, 244: 1275-1281 (1989); Verma, Scientific American: 68-84 (1990); and Miller, Nature, 357: 455-460 (1992). Introduction of any one of the nucleotides of the present invention or a gene encoding the polypeptides of the present invention can also be accomplished with extrachromosomal substrates (transient expression) or artificial chromosomes (stable expression). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes. Alternatively, it is contemplated that in other human disease states, preventing the expression of or inhibiting the activity of polypeptides of the invention will be useful in treating the disease states. It is contemplated that antisense therapy or gene therapy could be applied to negatively regulate the expression of polypeptides of the invention.

Other methods inhibiting expression of a protein include the introduction of antisense molecules to the nucleic acids of the present invention, their complements, or their translated RNA sequences, by methods known in the art. Further, the polypeptides of the present invention can be

inhibited by using targeted deletion methods, or the insertion of a negative regulatory element such as a silencer, which is tissue specific.

The present invention still further provides cells genetically engineered *in vivo* to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell. These methods can be used to increase or decrease the expression of the polynucleotides of the present invention.

Knowledge of DNA sequences provided by the invention allows for modification of cells to permit, increase, or decrease, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous promoter so that the cells express the protein at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the desired protein encoding sequences. See, for example, PCT International Publication No. WO 94/12650, PCT International Publication No. WO 92/20808, and PCT International Publication No. WO 91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the desired protein coding sequence, amplification of the marker DNA by standard selection methods results in co-amplification of the desired protein coding sequences in the cells.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequences include polyadenylation signals, mRNA stability elements, splice sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both

upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

4.9 TRANSGENIC ANIMALS

In preferred methods to determine biological functions of the polypeptides of the invention in vivo, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

Transgenic animals can be prepared wherein all or part of a promoter of the polynucleotides of the invention is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

The polynucleotides of the present invention also make possible the development, through, e.g., homologous recombination or knock out strategies, of animals that fail to express polypeptides of the invention or that express a variant polypeptide. Such animals are useful as models for studying the *in vivo* activities of polypeptide as well as for studying modulators of the polypeptides of the invention.

In preferred methods to determine biological functions of the polypeptides of the invention *in vivo*, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

Transgenic animals can be prepared wherein all or part of the polynucleotides of the invention promoter is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

4.10 USES AND BIOLOGICAL ACTIVITY

The polynucleotides and proteins of the present invention are expected to exhibit one or more of the uses or biological activities (including those associated with assays cited herein) identified herein. Uses or activities described for proteins of the present invention may be provided by administration or use of such proteins or of polynucleotides encoding such proteins (such as, for example, in gene therapies or vectors suitable for introduction of DNA). The mechanism underlying the particular condition or pathology will dictate whether the polypeptides of the invention, the polynucleotides of the invention or modulators (activators or inhibitors) thereof would be beneficial to the subject in need of treatment. Thus, "therapeutic compositions of the invention" include compositions comprising isolated polynucleotides (including recombinant DNA molecules, cloned genes and degenerate variants thereof) or polypeptides of the invention (including full length protein, mature protein and truncations or domains thereof), or compounds and other substances that modulate the overall activity of the target gene products, either at the level of target gene/protein expression or target protein activity. Such modulators include polypeptides, analogs, (variants), including fragments and fusion proteins, antibodies and other binding proteins; chemical compounds that directly or indirectly activate or inhibit the polypeptides of the invention (identified, e.g., via drug screening assays as described herein); antisense polynucleotides and polynucleotides suitable for triple helix formation; and in particular antibodies or other binding partners that specifically recognize one or more epitopes of the polypeptides of the invention.

The polypeptides of the present invention may likewise be involved in cellular activation or in one of the other physiological pathways described herein.

4.10.1 RESEARCH USES AND UTILITIES

The polynucleotides provided by the present invention can be used by the research community for various purposes. The polynucleotides can be used to express recombinant protein for analysis, characterization or therapeutic use; as markers for tissues in which the corresponding protein is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in disease states); as molecular weight markers on gels; as chromosome markers or tags (when labeled) to identify chromosomes or to map related gene positions; to compare with endogenous DNA sequences in patients to identify potential genetic disorders; as probes to hybridize and thus discover novel, related DNA sequences; as a source of information to derive PCR primers for genetic fingerprinting; as a probe to "subtract-out" known sequences in the process of discovering other novel polynucleotides; for selecting and making oligomers for attachment to a "gene chip" or other support, including for examination of expression patterns; to raise anti-protein antibodies using DNA immunization techniques; and as

an antigen to raise anti-DNA antibodies or elicit another immune response. Where the polynucleotide encodes a protein which binds or potentially binds to another protein (such as, for example, in a receptor-ligand interaction), the polynucleotide can also be used in interaction trap assays (such as, for example, that described in Gyuris et al., Cell 75:791-803 (1993)) to identify polynucleotides encoding the other protein with which binding occurs or to identify inhibitors of the binding interaction.

The polypeptides provided by the present invention can similarly be used in assays to determine biological activity, including in a panel of multiple proteins for high-throughput screening; to raise antibodies or to elicit another immune response; as a reagent (including the labeled reagent) in assays designed to quantitatively determine levels of the protein (or its receptor) in biological fluids; as markers for tissues in which the corresponding polypeptide is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in a disease state); and, of course, to isolate correlative receptors or ligands. Proteins involved in these binding interactions can also be used to screen for peptide or small molecule inhibitors or agonists of the binding interaction.

Any or all of these research utilities are capable of being developed into reagent grade or kit format for commercialization as research products.

Methods for performing the uses listed above are well known to those skilled in the art. References disclosing such methods include without limitation "Molecular Cloning: A Laboratory Manual", 2d ed., Cold Spring Harbor Laboratory Press, Sambrook, J., E. F. Fritsch and T. Maniatis eds., 1989, and "Methods in Enzymology: Guide to Molecular Cloning Techniques", Academic Press, Berger, S. L. and A. R. Kimmel eds., 1987.

4.10.2 NUTRITIONAL USES

Polynucleotides and polypeptides of the present invention can also be used as nutritional sources or supplements. Such uses include without limitation use as a protein or amino acid supplement, use as a carbon source, use as a nitrogen source and use as a source of carbohydrate. In such cases the polypeptide or polynucleotide of the invention can be added to the feed of a particular organism or can be administered as a separate solid or liquid preparation, such as in the form of powder, pills, solutions, suspensions or capsules. In the case of microorganisms, the polypeptide or polynucleotide of the invention can be added to the medium in or on which the microorganism is cultured.

4.10.3 CYTOKINE AND CELL PROLIFERATION/DIFFERENTIATION ACTIVITY

A polypeptide of the present invention may exhibit activity relating to cytokine, cell proliferation (either inducing or inhibiting) or cell differentiation (either inducing or inhibiting) activity or may induce production of other cytokines in certain cell populations. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Many protein factors discovered to date, including all known cytokines, have exhibited activity in one or more factor-dependent cell proliferation assays, and hence the assays serve as a convenient confirmation of cytokine activity. The activity of therapeutic compositions of the present invention is evidenced by any one of a number of routine factor dependent cell proliferation assays for cell lines including, without limitation, 32D, DA2, DA1G, T10, B9, B9/11, BaF3, MC9/G, M+(preB M+), 2E8, RB5, DA1, 123, T1165, HT2, CTLL2, TF-1, Mo7e, CMK, HUVEC, and Caco. Therapeutic compositions of the invention can be used in the following:

Assays for T-cell or thymocyte proliferation include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, *In Vitro* assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Bertagnolli et al., J. Immunol. 145:1706-1712, 1990; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Bertagnolli, et al., I. Immunol. 149:3778-3783, 1992; Bowman et al., I. Immunol. 152:1756-1761, 1994.

Assays for cytokine production and/or proliferation of spleen cells, lymph node cells or thymocytes include, without limitation, those described in: Polyclonal T cell stimulation, Kruisbeek, A. M. and Shevach, E. M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.12.1-3.12.14, John Wiley and Sons, Toronto. 1994; and Measurement of mouse and human interleukin-γ, Schreiber, R. D. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 6.8.1-6.8.8, John Wiley and Sons, Toronto. 1994.

Assays for proliferation and differentiation of hematopoietic and lymphopoietic cells include, without limitation, those described in: Measurement of Human and Murine Interleukin 2 and Interleukin 4, Bottomly, K., Davis, L. S. and Lipsky, P. E. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 6.3.1-6.3.12, John Wiley and Sons, Toronto. 1991; deVries et al., J. Exp. Med. 173:1205-1211, 1991; Moreau et al., Nature 336:690-692, 1988; Greenberger et al., Proc. Natl. Acad. Sci. U.S.A. 80:2931-2938, 1983; Measurement of mouse and human interleukin 6--Nordan, R. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.6.1-6.6.5, John Wiley and Sons, Toronto. 1991; Smith et al., Proc. Natl. Aced. Sci. U.S.A. 83:1857-1861, 1986; Measurement of human Interleukin 11--Bennett, F., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.15.1 John Wiley and Sons, Toronto. 1991; Measurement of mouse and human Interleukin

9--Ciarletta, A., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.13.1, John Wiley and Sons, Toronto. 1991.

Assays for T-cell clone responses to antigens (which will identify, among others, proteins that affect APC-T cell interactions as well as direct T-cell effects by measuring proliferation and cytokine production) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, *In Vitro* assays for Mouse Lymphocyte Function; Chapter 6, Cytokines and their cellular receptors; Chapter 7, Immunologic studies in Humans); Weinberger et al., Proc. Natl. Acad. Sci. USA 77:6091-6095, 1980; Weinberger et al., Eur. J. Immun. 11:405-411, 1981; Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988.

4.10.4 STEM CELL GROWTH FACTOR ACTIVITY

A polypeptide of the present invention may exhibit stem cell growth factor activity and be involved in the proliferation, differentiation and survival of pluripotent and totipotent stem cells including primordial germ cells, embryonic stem cells, hematopoietic stem cells and/or germ line stem cells. Administration of the polypeptide of the invention to stem cells *in vivo* or *ex vivo* is expected to maintain and expand cell populations in a totipotential or pluripotential state which would be useful for re-engineering damaged or diseased tissues, transplantation, manufacture of bio-pharmaceuticals and the development of bio-sensors. The ability to produce large quantities of human cells has important working applications for the production of human proteins which currently must be obtained from non-human sources or donors, implantation of cells to treat diseases such as Parkinson's, Alzheimer's and other neurodegenerative diseases; tissues for grafting such as bone marrow, skin, cartilage, tendons, bone, muscle (including cardiac muscle), blood vessels, cornea, neural cells, gastrointestinal cells and others; and organs for transplantation such as kidney, liver, pancreas (including islet cells), heart and lung.

It is contemplated that multiple different exogenous growth factors and/or cytokines may be administered in combination with the polypeptide of the invention to achieve the desired effect, including any of the growth factors listed herein, other stem cell maintenance factors, and specifically including stem cell factor (SCF), leukemia inhibitory factor (LIF), Flt-3 ligand (Flt-3L), any of the interleukins, recombinant soluble IL-6 receptor fused to IL-6, macrophage inflammatory protein 1-alpha (MIP-1-alpha), G-CSF, GM-CSF, thrombopoietin (TPO), platelet factor 4 (PF-4), platelet-derived growth factor (PDGF), neural growth factors and basic fibroblast growth factor (bFGF).

Since totipotent stem cells can give rise to virtually any mature cell type, expansion of these cells in culture will facilitate the production of large quantities of mature cells. Techniques for culturing stem cells are known in the art and administration of polypeptides of the invention, optionally with other growth factors and/or cytokines, is expected to enhance the survival and proliferation of the stem cell populations. This can be accomplished by direct administration of the polypeptide of the invention to the culture medium. Alternatively, stroma cells transfected with a polynucleotide that encodes for the polypeptide of the invention can be used as a feeder layer for the stem cell populations in culture or in vivo. Stromal support cells for feeder layers may include embryonic bone marrow fibroblasts, bone marrow stromal cells, fetal liver cells, or cultured embryonic fibroblasts (see U.S. Patent No. 5,690,926).

Stem cells themselves can be transfected with a polynucleotide of the invention to induce autocrine expression of the polypeptide of the invention. This will allow for generation of undifferentiated totipotential/pluripotential stem cell lines that are useful as is or that can then be differentiated into the desired mature cell types. These stable cell lines can also serve as a source of undifferentiated totipotential/pluripotential mRNA to create cDNA libraries and templates for polymerase chain reaction experiments. These studies would allow for the isolation and identification of differentially expressed genes in stem cell populations that regulate stem cell proliferation and/or maintenance.

Expansion and maintenance of totipotent stem cell populations will be useful in the treatment of many pathological conditions. For example, polypeptides of the present invention may be used to manipulate stem cells in culture to give rise to neuroepithelial cells that can be used to augment or replace cells damaged by illness, autoimmune disease, accidental damage or genetic disorders. The polypeptide of the invention may be useful for inducing the proliferation of neural cells and for the regeneration of nerve and brain tissue, i.e. for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders which involve degeneration, death or trauma to neural cells or nerve tissue. In addition, the expanded stem cell populations can also be genetically altered for gene therapy purposes and to decrease host rejection of replacement tissues after grafting or implantation.

Expression of the polypeptide of the invention and its effect on stem cells can also be manipulated to achieve controlled differentiation of the stem cells into more differentiated cell types. A broadly applicable method of obtaining pure populations of a specific differentiated cell type from undifferentiated stem cell populations involves the use of a cell-type specific promoter driving a selectable marker. The selectable marker allows only cells of the desired type to survive. For example, stem cells can be induced to differentiate into cardiomyocytes (Wobus et al., Differentiation, 48: 173-182, (1991); Klug et al., J. Clin. Invest., 98(1): 216-224, (1998))

or skeletal muscle cells (Browder, L. W. In: *Principles of Tissue Engineering eds.* Lanza et al., Academic Press (1997)). Alternatively, directed differentiation of stem cells can be accomplished by culturing the stem cells in the presence of a differentiation factor such as retinoic acid and an antagonist of the polypeptide of the invention which would inhibit the effects of endogenous stem cell factor activity and allow differentiation to proceed.

In vitro cultures of stem cells can be used to determine if the polypeptide of the invention exhibits stem cell growth factor activity. Stem cells are isolated from any one of various cell sources (including hematopoietic stem cells and embryonic stem cells) and cultured on a feeder layer, as described by Thompson et al. Proc. Natl. Acad. Sci, U.S.A., 92: 7844-7848 (1995), in the presence of the polypeptide of the invention alone or in combination with other growth factors or cytokines. The ability of the polypeptide of the invention to induce stem cells proliferation is determined by colony formation on semi-solid support e.g. as described by Bernstein et al., Blood, 77: 2316-2321 (1991).

4.10.5 HEMATOPOIESIS REGULATING ACTIVITY

A polypeptide of the present invention may be involved in regulation of hematopoiesis and, consequently, in the treatment of myeloid or lymphoid cell disorders. Even marginal biological activity in support of colony forming cells or of factor-dependent cell lines indicates involvement in regulating hematopoiesis, e.g. in supporting the growth and proliferation of erythroid progenitor cells alone or in combination with other cytokines, thereby indicating utility, for example, in treating various anemias or for use in conjunction with irradiation/chemotherapy to stimulate the production of erythroid precursors and/or erythroid cells; in supporting the growth and proliferation of myeloid cells such as granulocytes and monocytes/macrophages (i.e., traditional CSF activity) useful, for example, in conjunction with chemotherapy to prevent or treat consequent myelo-suppression; in supporting the growth and proliferation of megakaryocytes and consequently of platelets thereby allowing prevention or treatment of various platelet disorders such as thrombocytopenia, and generally for use in place of or complimentary to platelet transfusions; and/or in supporting the growth and proliferation of hematopoietic stem cells which are capable of maturing to any and all of the above-mentioned hematopoietic cells and therefore find therapeutic utility in various stem cell disorders (such as those usually treated with transplantation, including, without limitation, aplastic anemia and paroxysmal nocturnal hemoglobinuria), as well as in repopulating the stem cell compartment post irradiation/chemotherapy, either in-vivo or ex-vivo (i.e., in conjunction with bone marrow transplantation or with peripheral progenitor cell transplantation (homologous or heterologous)) as normal cells or genetically manipulated for gene therapy.

Therapeutic compositions of the invention can be used in the following:

Suitable assays for proliferation and differentiation of various hematopoietic lines are cited above.

Assays for embryonic stem cell differentiation (which will identify, among others, proteins that influence embryonic differentiation hematopoiesis) include, without limitation, those described in: Johansson et al. Cellular Biology 15:141-151, 1995; Keller et al., Molecular and Cellular Biology 13:473-486, 1993; McClanahan et al., Blood 81:2903-2915, 1993.

Assays for stem cell survival and differentiation (which will identify, among others, proteins that regulate lympho-hematopoiesis) include, without limitation, those described in: Methylcellulose colony forming assays, Freshney, M. G. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 265-268, Wiley-Liss, Inc., New York, N.Y. 1994; Hirayama et al., Proc. Natl. Acad. Sci. USA 89:5907-5911, 1992; Primitive hematopoietic colony forming cells with high proliferative potential, McNiece, I. K. and Briddell, R. A. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 23-39, Wiley-Liss, Inc., New York, N.Y. 1994; Neben et al., Experimental Hematology 22:353-359, 1994; Cobblestone area forming cell assay, Ploemacher, R. E. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 1-21, Wiley-Liss, Inc., New York, N.Y. 1994; Long term bone marrow cultures in the presence of stromal cells, Spooncer, E., Dexter, M. and Allen, T. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 163-179, Wiley-Liss, Inc., New York, N.Y. 1994; Long term culture initiating cell assay, Sutherland, H. J. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 139-162, Wiley-Liss, Inc., New York, N.Y. 1994.

4.10.6 TISSUE GROWTH ACTIVITY

A polypeptide of the present invention also may be involved in bone, cartilage, tendon, ligament and/or nerve tissue growth or regeneration, as well as in wound healing and tissue repair and replacement, and in healing of burns, incisions and ulcers.

A polypeptide of the present invention which induces cartilage and/or bone growth in circumstances where bone is not normally formed, has application in the healing of bone fractures and cartilage damage or defects in humans and other animals. Compositions of a polypeptide, antibody, binding partner, or other modulator of the invention may have prophylactic use in closed as well as open fracture reduction and also in the improved fixation of artificial joints. De novo bone formation induced by an osteogenic agent contributes to the repair of congenital, trauma induced, or oncologic resection induced craniofacial defects, and also is useful in cosmetic plastic surgery.

A polypeptide of this invention may also be involved in attracting bone-forming cells, stimulating growth of bone-forming cells, or inducing differentiation of progenitors of bone-forming cells. Treatment of osteoporosis, osteoarthritis, bone degenerative disorders, or periodontal disease, such as through stimulation of bone and/or cartilage repair or by blocking inflammation or processes of tissue destruction (collagenase activity, osteoclast activity, etc.) mediated by inflammatory processes may also be possible using the composition of the invention.

Another category of tissue regeneration activity that may involve the polypeptide of the present invention is tendon/ligament formation. Induction of tendon/ligament-like tissue or other tissue formation in circumstances where such tissue is not normally formed, has application in the healing of tendon or ligament tears, deformities and other tendon or ligament defects in humans and other animals. Such a preparation employing a tendon/ligament-like tissue inducing protein may have prophylactic use in preventing damage to tendon or ligament tissue, as well as use in the improved fixation of tendon or ligament to bone or other tissues, and in repairing defects to tendon or ligament tissue. De novo tendon/ligament-like tissue formation induced by a composition of the present invention contributes to the repair of congenital, trauma induced, or other tendon or ligament defects of other origin, and is also useful in cosmetic plastic surgery for attachment or repair of tendons or ligaments. The compositions of the present invention may provide environment to attract tendon- or ligament-forming cells, stimulate growth of tendon- or ligament-forming cells, induce differentiation of progenitors of tendon- or ligament-forming cells, or induce growth of tendon/ligament cells or progenitors ex vivo for return in vivo to effect tissue repair. The compositions of the invention may also be useful in the treatment of tendinitis, carpal tunnel syndrome and other tendon or ligament defects. The compositions may also include an appropriate matrix and/or sequestering agent as a carrier as is well known in the art.

The compositions of the present invention may also be useful for proliferation of neural cells and for regeneration of nerve and brain tissue, i.e. for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders, which involve degeneration, death or trauma to neural cells or nerve tissue. More specifically, a composition may be used in the treatment of diseases of the peripheral nervous system, such as peripheral nerve injuries, peripheral neuropathy and localized neuropathies, and central nervous system diseases, such as Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome. Further conditions which may be treated in accordance with the present invention include mechanical and traumatic disorders, such as spinal cord disorders, head trauma and cerebrovascular diseases such as stroke. Peripheral neuropathies

resulting from chemotherapy or other medical therapies may also be treatable using a composition of the invention.

Compositions of the invention may also be useful to promote better or faster closure of non-healing wounds, including without limitation pressure ulcers, ulcers associated with vascular insufficiency, surgical and traumatic wounds, and the like.

Compositions of the present invention may also be involved in the generation or regeneration of other tissues, such as organs (including, for example, pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac) and vascular (including vascular endothelium) tissue, or for promoting the growth of cells comprising such tissues. Part of the desired effects may be by inhibition or modulation of fibrotic scarring may allow normal tissue to regenerate. A polypeptide of the present invention may also exhibit angiogenic activity.

A composition of the present invention may also be useful for gut protection or regeneration and treatment of lung or liver fibrosis, reperfusion injury in various tissues, and conditions resulting from systemic cytokine damage.

A composition of the present invention may also be useful for promoting or inhibiting differentiation of tissues described above from precursor tissues or cells; or for inhibiting the growth of tissues described above.

Therapeutic compositions of the invention can be used in the following:

Assays for tissue generation activity include, without limitation, those described in: International Patent Publication No. WO95/16035 (bone, cartilage, tendon); International Patent Publication No. WO95/05846 (nerve, neuronal); International Patent Publication No. WO91/07491 (skin, endothelium).

Assays for wound healing activity include, without limitation, those described in: Winter, Epidermal Wound Healing, pps. 71-112 (Maibach, H. I. and Rovee, D. T., eds.), Year Book Medical Publishers, Inc., Chicago, as modified by Eaglstein and Mertz, J. Invest. Dermatol 71:382-84 (1978).

4.10.7 IMMUNE STIMULATING OR SUPPRESSING ACTIVITY

A polypeptide of the present invention may also exhibit immune stimulating or immune suppressing activity, including without limitation the activities for which assays are described herein. A polynucleotide of the invention can encode a polypeptide exhibiting such activities. A protein may be useful in the treatment of various immune deficiencies and disorders (including severe combined immunodeficiency (SCID)), e.g., in regulating (up or down) growth and proliferation of T and/or B lymphocytes, as well as effecting the cytolytic activity of NK cells and other cell populations. These immune deficiencies may be genetic or be caused by viral (e.g.,

HIV) as well as bacterial or fungal infections, or may result from autoimmune disorders. More specifically, infectious diseases causes by viral, bacterial, fungal or other infection may be treatable using a protein of the present invention, including infections by HIV, hepatitis viruses, herpes viruses, mycobacteria, Leishmania spp., malaria spp. and various fungal infections such as candidiasis. Of course, in this regard, proteins of the present invention may also be useful where a boost to the immune system generally may be desirable, i.e., in the treatment of cancer.

Autoimmune disorders which may be treated using a protein of the present invention include, for example, connective tissue disease, multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, autoimmune pulmonary inflammation, Guillain-Barre syndrome, autoimmune thyroiditis, insulin dependent diabetes mellitis, myasthenia gravis, graft-versus-host disease and autoimmune inflammatory eye disease. Such a protein (or antagonists thereof, including antibodies) of the present invention may also to be useful in the treatment of allergic reactions and conditions (e.g., anaphylaxis, serum sickness, drug reactions, food allergies, insect venom allergies, mastocytosis, allergic rhinitis, hypersensitivity pneumonitis, urticaria, angioedema, eczema, atopic dermatitis, allergic contact dermatitis, erythema multiforme, Stevens-Johnson syndrome, allergic conjunctivitis, atopic keratoconjunctivitis, venereal keratoconjunctivitis, giant papillary conjunctivitis and contact allergies), such as asthma (particularly allergic asthma) or other respiratory problems. Other conditions, in which immune suppression is desired (including, for example, organ transplantation), may also be treatable using a protein (or antagonists thereof) of the present invention. The therapeutic effects of the polypeptides or antagonists thereof on allergic reactions can be evaluated by in vivo animals models such as the cumulative contact enhancement test (Lastborn et al., Toxicology 125: 59-66, 1998), skin prick test (Hoffmann et al., Allergy 54: 446-54, 1999), guinea pig skin sensitization test (Vohr et al., Arch. Toxocol. 73: 501-9), and murine local lymph node assay (Kimber et al., J. Toxicol. Environ. Health 53: 563-79).

Using the proteins of the invention it may also be possible to modulate immune responses, in a number of ways. Down regulation may be in the form of inhibiting or blocking an immune response already in progress or may involve preventing the induction of an immune response. The functions of activated T cells may be inhibited by suppressing T cell responses or by inducing specific tolerance in T cells, or both. Immunosuppression of T cell responses is generally an active, non-antigen-specific, process which requires continuous exposure of the T cells to the suppressive agent. Tolerance, which involves inducing non-responsiveness or anergy in T cells, is distinguishable from immunosuppression in that it is generally antigen-specific and persists after exposure to the tolerizing agent has ceased. Operationally, tolerance can be

demonstrated by the lack of a T cell response upon reexposure to specific antigen in the absence of the tolerizing agent.

Down regulating or preventing one or more antigen functions (including without limitation B lymphocyte antigen functions (such as, for example, B7)), e.g., preventing high level lymphokine synthesis by activated T cells, will be useful in situations of tissue, skin and organ transplantation and in graft-versus-host disease (GVHD). For example, blockage of T cell function should result in reduced tissue destruction in tissue transplantation. Typically, in tissue transplants, rejection of the transplant is initiated through its recognition as foreign by T cells, followed by an immune reaction that destroys the transplant. The administration of a therapeutic composition of the invention may prevent cytokine synthesis by immune cells, such as T cells, and thus acts as an immunosuppressant. Moreover, a lack of costimulation may also be sufficient to anergize the T cells, thereby inducing tolerance in a subject. Induction of long-term tolerance by B lymphocyte antigen-blocking reagents may avoid the necessity of repeated administration of these blocking reagents. To achieve sufficient immunosuppression or tolerance in a subject, it may also be necessary to block the function of a combination of B lymphocyte antigens.

The efficacy of particular therapeutic compositions in preventing organ transplant rejection or GVHD can be assessed using animal models that are predictive of efficacy in humans. Examples of appropriate systems which can be used include allogeneic cardiac grafts in rats and xenogeneic pancreatic islet cell grafts in mice, both of which have been used to examine the immunosuppressive effects of CTLA4Ig fusion proteins in vivo as described in Lenschow et al., Science 257:789-792 (1992) and Turka et al., Proc. Natl. Acad. Sci USA, 89:11102-11105 (1992). In addition, murine models of GVHD (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 846-847) can be used to determine the effect of therapeutic compositions of the invention on the development of that disease.

Blocking antigen function may also be therapeutically useful for treating autoimmune diseases. Many autoimmune disorders are the result of inappropriate activation of T cells that are reactive against self tissue and which promote the production of cytokines and autoantibodies involved in the pathology of the diseases. Preventing the activation of autoreactive T cells may reduce or eliminate disease symptoms. Administration of reagents which block stimulation of T cells can be used to inhibit T cell activation and prevent production of autoantibodies or T cell-derived cytokines which may be involved in the disease process. Additionally, blocking reagents may induce antigen-specific tolerance of autoreactive T cells which could lead to long-term relief from the disease. The efficacy of blocking reagents in preventing or alleviating autoimmune disorders can be determined using a number of well-characterized animal models of human autoimmune diseases. Examples include murine experimental autoimmune encephalitis,

systemic lupus erythmatosis in MRL/lpr/lpr mice or NZB hybrid mice, murine autoimmune collagen arthritis, diabetes mellitus in NOD mice and BB rats, and murine experimental myasthenia gravis (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 840-856).

Upregulation of an antigen function (e.g., a B lymphocyte antigen function), as a means of up regulating immune responses, may also be useful in therapy. Upregulation of immune responses may be in the form of enhancing an existing immune response or eliciting an initial immune response. For example, enhancing an immune response may be useful in cases of viral infection, including systemic viral diseases such as influenza, the common cold, and encephalitis.

Alternatively, anti-viral immune responses may be enhanced in an infected patient by removing T cells from the patient, costimulating the T cells in vitro with viral antigen-pulsed APCs either expressing a peptide of the present invention or together with a stimulatory form of a soluble peptide of the present invention and reintroducing the in vitro activated T cells into the patient. Another method of enhancing anti-viral immune responses would be to isolate infected cells from a patient, transfect them with a nucleic acid encoding a protein of the present invention as described herein such that the cells express all or a portion of the protein on their surface, and reintroduce the transfected cells into the patient. The infected cells would now be capable of delivering a costimulatory signal to, and thereby activate, T cells in vivo.

A polypeptide of the present invention may provide the necessary stimulation signal to T cells to induce a T cell mediated immune response against the transfected tumor cells. In addition, tumor cells which lack MHC class I or MHC class II molecules, or which fail to reexpress sufficient mounts of MHC class I or MHC class II molecules, can be transfected with nucleic acid encoding all or a portion of (e.g., a cytoplasmic-domain truncated portion) of an MHC class I alpha chain protein and β_2 microglobulin protein or an MHC class II alpha chain protein and an MHC class II beta chain protein to thereby express MHC class I or MHC class II proteins on the cell surface. Expression of the appropriate class I or class II MHC in conjunction with a peptide having the activity of a B lymphocyte antigen (e.g., B7-1, B7-2, B7-3) induces a T cell mediated immune response against the transfected tumor cell. Optionally, a gene encoding an antisense construct which blocks expression of an MHC class II associated protein, such as the invariant chain, can also be cotransfected with a DNA encoding a peptide having the activity of a B lymphocyte antigen to promote presentation of tumor associated antigens and induce tumor specific immunity. Thus, the induction of a T cell mediated immune response in a human subject may be sufficient to overcome tumor-specific tolerance in the subject.

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The activity of a protein of the invention may, among other means, be measured by the following methods:

Suitable assays for thymocyte or splenocyte cytotoxicity include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Herrmann et al., Proc. Natl. Acad. Sci. USA 78:2488-2492, 1981; Herrmann et al., J. Immunol. 128:1968-1974, 1982; Handa et al., J. Immunol. 135:1564-1572, 1985; Takai et al., I. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bowman et al., J. Virology 61:1992-1998; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Brown et al., J. Immunol. 153:3079-3092, 1994.

Assays for T-cell-dependent immunoglobulin responses and isotype switching (which will identify, among others, proteins that modulate T-cell dependent antibody responses and that affect Th1/Th2 profiles) include, without limitation, those described in: Maliszewski, J. Immunol. 144:3028-3033, 1990; and Assays for B cell function: In vitro antibody production, Mond, J. J. and Brunswick, M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.8.1-3.8.16, John Wiley and Sons, Toronto. 1994.

Mixed lymphocyte reaction (MLR) assays (which will identify, among others, proteins that generate predominantly Th1 and CTL responses) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bertagnolli et al., J. Immunol. 149:3778-3783, 1992.

Dendritic cell-dependent assays (which will identify, among others, proteins expressed by dendritic cells that activate naive T-cells) include, without limitation, those described in: Guery et al., J. Immunol, 134:536-544, 1995; Inaba et al., Journal of Experimental Medicine 173:549-559, 1991; Macatonia et al., Journal of Immunology 154:5071-5079, 1995; Porgador et al., Journal of Experimental Medicine 182:255-260, 1995; Nair et al., Journal of Virology 67:4062-4069, 1993; Huang et al., Science 264:961-965, 1994; Macatonia et al., Journal of Experimental Medicine 169:1255-1264, 1989; Bhardwaj et al., Journal of Clinical Investigation 94:797-807, 1994; and Inaba et al., Journal of Experimental Medicine 172:631-640, 1990.

Assays for lymphocyte survival/apoptosis (which will identify, among others, proteins that prevent apoptosis after superantigen induction and proteins that regulate lymphocyte homeostasis) include, without limitation, those described in: Darzynkiewicz et al., Cytometry

13:795-808, 1992; Gorczyca et al., Leukemia 7:659-670, 1993; Gorczyca et al., Cancer Research 53:1945-1951, 1993; Itoh et al., Cell 66:233-243, 1991; Zacharchuk, Journal of Immunology 145:4037-4045, 1990; Zamai et al., Cytometry 14:891-897, 1993; Gorczyca et al., International Journal of Oncology 1:639-648, 1992.

Assays for proteins that influence early steps of T-cell commitment and development include, without limitation, those described in: Antica et al., Blood 84:111-117, 1994; Fine et al., Cellular Immunology 155:111-122, 1994; Galy et al., Blood 85:2770-2778, 1995; Toki et al., Proc. Nat. Acad Sci. USA 88:7548-7551, 1991.

4.10.8 ACTIVIN/INHIBIN ACTIVITY

A polypeptide of the present invention may also exhibit activin- or inhibin-related activities. A polynucleotide of the invention may encode a polypeptide exhibiting such characteristics. Inhibins are characterized by their ability to inhibit the release of follicle stimulating hormone (FSH), while activins and are characterized by their ability to stimulate the release of follicle stimulating hormone (FSH). Thus, a polypeptide of the present invention, alone or in heterodimers with a member of the inhibin family, may be useful as a contraceptive based on the ability of inhibins to decrease fertility in female mammals and decrease spermatogenesis in male mammals. Administration of sufficient amounts of other inhibins can induce infertility in these mammals. Alternatively, the polypeptide of the invention, as a homodimer or as a heterodimer with other protein subunits of the inhibin group, may be useful as a fertility inducing therapeutic, based upon the ability of activin molecules in stimulating FSH release from cells of the anterior pituitary. See, for example, U.S. Pat. No. 4,798,885. A polypeptide of the invention may also be useful for advancement of the onset of fertility in sexually immature mammals, so as to increase the lifetime reproductive performance of domestic animals such as, but not limited to, cows, sheep and pigs.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods.

Assays for activin/inhibin activity include, without limitation, those described in: Vale et al., Endocrinology 91:562-572, 1972; Ling et al., Nature 321:779-782, 1986; Vale et al., Nature 321:776-779, 1986; Mason et al., Nature 318:659-663, 1985; Forage et al., Proc. Natl. Acad. Sci. USA 83:3091-3095, 1986.

4.10.9 CHEMOTACTIC/CHEMOKINETIC ACTIVITY

A polypeptide of the present invention may be involved in chemotactic or chemokinetic activity for mammalian cells, including, for example, monocytes, fibroblasts, neutrophils,

T-cells, mast cells, eosinophils, epithelial and/or endothelial cells. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Chemotactic and chemokinetic receptor activation can be used to mobilize or attract a desired cell population to a desired site of action. Chemotactic or chemokinetic compositions (e.g. proteins, antibodies, binding partners, or modulators of the invention) provide particular advantages in treatment of wounds and other trauma to tissues, as well as in treatment of localized infections. For example, attraction of lymphocytes, monocytes or neutrophils to tumors or sites of infection may result in improved immune responses against the tumor or infecting agent.

A protein or peptide has chemotactic activity for a particular cell population if it can stimulate, directly or indirectly, the directed orientation or movement of such cell population. Preferably, the protein or peptide has the ability to directly stimulate directed movement of cells. Whether a particular protein has chemotactic activity for a population of cells can be readily determined by employing such protein or peptide in any known assay for cell chemotaxis.

Therapeutic compositions of the invention can be used in the following:

Assays for chemotactic activity (which will identify proteins that induce or prevent chemotaxis) consist of assays that measure the ability of a protein to induce the migration of cells across a membrane as well as the ability of a protein to induce the adhesion of one cell population to another cell population. Suitable assays for movement and adhesion include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Marguiles, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 6.12, Measurement of alpha and beta Chemokines 6.12.1-6.12.28; Taub et al. J. Clin. Invest. 95:1370-1376, 1995; Lind et al. APMIS 103:140-146, 1995; Muller et al Eur. J. Immunol. 25:1744-1748; Gruber et al. J. of Immunol. 152:5860-5867, 1994; Johnston et al. J. of Immunol. 153:1762-1768, 1994.

4.10.10 HEMOSTATIC AND THROMBOLYTIC ACTIVITY

A polypeptide of the invention may also be involved in hemostasis or thrombolysis or thrombosis. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Compositions may be useful in treatment of various coagulation disorders (including hereditary disorders, such as hemophilias) or to enhance coagulation and other hemostatic events in treating wounds resulting from trauma, surgery or other causes. A composition of the invention may also be useful for dissolving or inhibiting formation of thromboses and for treatment and prevention of conditions resulting therefrom (such as, for example, infarction of cardiac and central nervous system vessels (e.g., stroke).

Therapeutic compositions of the invention can be used in the following:

Assay for hemostatic and thrombolytic activity include, without limitation, those described in: Linet et al., J. Clin. Pharmacol. 26:131-140, 1986; Burdick et al., Thrombosis Res. 45:413-419, 1987; Humphrey et al., Fibrinolysis 5:71-79 (1991); Schaub, Prostaglandins 35:467-474, 1988.

4.10.11 CANCER DIAGNOSIS AND THERAPY

Polypeptides of the invention may be involved in cancer cell generation, proliferation or metastasis. Detection of the presence or amount of polynucleotides or polypeptides of the invention may be useful for the diagnosis and/or prognosis of one or more types of cancer. For example, the presence or increased expression of a polynucleotide/polypeptide of the invention may indicate a hereditary risk of cancer, a precancerous condition, or an ongoing malignancy. Conversely, a defect in the gene or absence of the polypeptide may be associated with a cancer condition. Identification of single nucleotide polymorphisms associated with cancer or a predisposition to cancer may also be useful for diagnosis or prognosis.

Cancer treatments promote tumor regression by inhibiting tumor cell proliferation, inhibiting angiogenesis (growth of new blood vessels that is necessary to support tumor growth) and/or prohibiting metastasis by reducing tumor cell motility or invasiveness. Therapeutic compositions of the invention may be effective in adult and pediatric oncology including in solid phase tumors/malignancies, locally advanced tumors, human soft tissue sarcomas, metastatic cancer, including lymphatic metastases, blood cell malignancies including multiple myeloma, acute and chronic leukemias, and lymphomas, head and neck cancers including mouth cancer, larynx cancer and thyroid cancer, lung cancers including small cell carcinoma and non-small cell cancers, breast cancers including small cell carcinoma and ductal carcinoma, gastrointestinal cancers including esophageal cancer, stomach cancer, colon cancer, colorectal cancer and polyps associated with colorectal neoplasia, pancreatic cancers, liver cancer, urologic cancers including bladder cancer and prostate cancer, malignancies of the female genital tract including ovarian carcinoma, uterine (including endometrial) cancers, and solid tumor in the ovarian follicle, kidney cancers including renal cell carcinoma, brain cancers including intrinsic brain tumors, neuroblastoma, astrocytic brain tumors, gliomas, metastatic tumor cell invasion in the central nervous system, bone cancers including osteomas, skin cancers including malignant melanoma, tumor progression of human skin keratinocytes, squamous cell carcinoma, basal cell carcinoma, hemangiopericytoma and Karposi's sarcoma.

Polypeptides, polynucleotides, or modulators of polypeptides of the invention (including inhibitors and stimulators of the biological activity of the polypeptide of the invention) may be administered to treat cancer. Therapeutic compositions can be administered in therapeutically

effective dosages alone or in combination with adjuvant cancer therapy such as surgery, chemotherapy, radiotherapy, thermotherapy, and laser therapy, and may provide a beneficial effect, e.g. reducing tumor size, slowing rate of tumor growth, inhibiting metastasis, or otherwise improving overall clinical condition, without necessarily eradicating the cancer.

The composition can also be administered in therapeutically effective amounts as a portion of an anti-cancer cocktail. An anti-cancer cocktail is a mixture of the polypeptide or modulator of the invention with one or more anti-cancer drugs in addition to a pharmaceutically acceptable carrier for delivery. The use of anti-cancer cocktails as a cancer treatment is routine. Anti-cancer drugs that are well known in the art and can be used as a treatment in combination with the polypeptide or modulator of the invention include: Actinomycin D, Aminoglutethimide, Asparaginase, Bleomycin, Busulfan, Carboplatin, Carmustine, Chlorambucil, Cisplatin (cis-DDP), Cyclophosphamide, Cytarabine HCl (Cytosine arabinoside), Dacarbazine, Dactinomycin, Daunorubicin HCl, Doxorubicin HCl, Estramustine phosphate sodium, Etoposide (V16-213), Floxuridine, 5-Fluorouracil (5-Fu), Flutamide, Hydroxyurea (hydroxycarbamide), Ifosfamide, Interferon Alpha-2a, Interferon Alpha-2b, Leuprolide acetate (LHRH-releasing factor analog), Lomustine, Mechlorethamine HCl (nitrogen mustard), Melphalan, Mercaptopurine, Mesna, Methotrexate (MTX), Mitomycin, Mitoxantrone HCl, Octreotide, Plicamycin, Procarbazine HCl, Streptozocin, Tamoxifen citrate, Thioguanine, Thiotepa, Vinblastine sulfate, Vincristine sulfate, Amsacrine, Azacitidine, Hexamethylmelamine, Interleukin-2, Mitoguazone, Pentostatin, Semustine, Teniposide, and Vindesine sulfate.

In addition, therapeutic compositions of the invention may be used for prophylactic treatment of cancer. There are hereditary conditions and/or environmental situations (e.g. exposure to carcinogens) known in the art that predispose an individual to developing cancers. Under these circumstances, it may be beneficial to treat these individuals with therapeutically effective doses of the polypeptide of the invention to reduce the risk of developing cancers.

In vitro models can be used to determine the effective doses of the polypeptide of the invention as a potential cancer treatment. These in vitro models include proliferation assays of cultured tumor cells, growth of cultured tumor cells in soft agar (see Freshney, (1987) Culture of Animal Cells: A Manual of Basic Technique, Wily-Liss, New York, NY Ch 18 and Ch 21), tumor systems in nude mice as described in Giovanella et al., J. Natl. Can. Inst., 52: 921-30 (1974), mobility and invasive potential of tumor cells in Boyden Chamber assays as described in Pilkington et al., Anticancer Res., 17: 4107-9 (1997), and angiogenesis assays such as induction of vascularization of the chick chorioallantoic membrane or induction of vascular endothelial cell migration as described in Ribatta et al., Intl. J. Dev. Biol., 40: 1189-97 (1999) and Li et al., Clin. Exp. Metastasis, 17:423-9 (1999), respectively. Suitable tumor cells lines are available,

e.g. from American Type Tissue Culture Collection catalogs.

4.10.12 RECEPTOR/LIGAND ACTIVITY

A polypeptide of the present invention may also demonstrate activity as receptor, receptor ligand or inhibitor or agonist of receptor/ligand interactions. A polynucleotide of the invention can encode a polypeptide exhibiting such characteristics. Examples of such receptors and ligands include, without limitation, cytokine receptors and their ligands, receptor kinases and their ligands, receptor phosphatases and their ligands, receptors involved in cell-cell interactions and their ligands (including without limitation, cellular adhesion molecules (such as selectins, integrins and their ligands) and receptor/ligand pairs involved in antigen presentation, antigen recognition and development of cellular and humoral immune responses. Receptors and ligands are also useful for screening of potential peptide or small molecule inhibitors of the relevant receptor/ligand interaction. A protein of the present invention (including, without limitation, fragments of receptors and ligands) may themselves be useful as inhibitors of receptor/ligand interactions.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods:

Suitable assays for receptor-ligand activity include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley- Interscience (Chapter 7.28, Measurement of Cellular Adhesion under static conditions 7.28.1-7.28.22), Takai et al., Proc. Natl. Acad. Sci. USA 84:6864-6868, 1987; Bierer et al., J. Exp. Med. 168:1145-1156, 1988; Rosenstein et al., J. Exp. Med. 169:149-160 1989; Stoltenborg et al., J. Immunol. Methods 175:59-68, 1994; Stitt et al., Cell 80:661-670, 1995.

By way of example, the polypeptides of the invention may be used as a receptor for a ligand(s) thereby transmitting the biological activity of that ligand(s). Ligands may be identified through binding assays, affinity chromatography, dihybrid screening assays, BIAcore assays, gel overlay assays, or other methods known in the art.

Studies characterizing drugs or proteins as agonist or antagonist or partial agonists or a partial antagonist require the use of other proteins as competing ligands. The polypeptides of the present invention or ligand(s) thereof may be labeled by being coupled to radioisotopes, colorimetric molecules or a toxin molecules by conventional methods. ("Guide to Protein Purification" Murray P. Deutscher (ed) Methods in Enzymology Vol. 182 (1990) Academic Press, Inc. San Diego). Examples of radioisotopes include, but are not limited to, tritium and carbon-14. Examples of colorimetric molecules include, but are not limited to, fluorescent

molecules such as fluorescamine, or rhodamine or other colorimetric molecules. Examples of toxins include, but are not limited, to ricin.

4.10.13 DRUG SCREENING

This invention is particularly useful for screening chemical compounds by using the novel polypeptides or binding fragments thereof in any of a variety of drug screening techniques. The polypeptides or fragments employed in such a test may either be free in solution, affixed to a solid support, borne on a cell surface or located intracellularly. One method of drug screening utilizes eukaryotic or prokaryotic host cells which are stably transformed with recombinant nucleic acids expressing the polypeptide or a fragment thereof. Drugs are screened against such transformed cells in competitive binding assays. Such cells, either in viable or fixed form, can be used for standard binding assays. One may measure, for example, the formation of complexes between polypeptides of the invention or fragments and the agent being tested or examine the diminution in complex formation between the novel polypeptides and an appropriate cell line, which are well known in the art.

Sources for test compounds that may be screened for ability to bind to or modulate (i.e., increase or decrease) the activity of polypeptides of the invention include (1) inorganic and organic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of either random or mimetic peptides, oligonucleotides or organic molecules.

Chemical libraries may be readily synthesized or purchased from a number of commercial sources, and may include structural analogs of known compounds or compounds that are identified as "hits" or "leads" via natural product screening.

The sources of natural product libraries are microorganisms (including bacteria and fungi), animals, plants or other vegetation, or marine organisms, and libraries of mixtures for screening may be created by: (1) fermentation and extraction of broths from soil, plant or marine microorganisms or (2) extraction of the organisms themselves. Natural product libraries include polyketides, non-ribosomal peptides, and (non-naturally occurring) variants thereof. For a review, see *Science 282*:63-68 (1998).

Combinatorial libraries are composed of large numbers of peptides, oligonucleotides or organic compounds and can be readily prepared by traditional automated synthesis methods, PCR, cloning or proprietary synthetic methods. Of particular interest are peptide and oligonucleotide combinatorial libraries. Still other libraries of interest include peptide, protein, peptidomimetic, multiparallel synthetic collection, recombinatorial, and polypeptide libraries. For a review of combinatorial chemistry and libraries created therefrom, see Myers, Curr. Opin. Biotechnol. 8:701-707 (1997). For reviews and examples of peptidomimetic libraries, see

Al-Obeidi et al., Mol. Biotechnol, 9(3):205-23 (1998); Hruby et al., Curr Opin Chem Biol, 1(1):114-19 (1997); Dorner et al., Bioorg Med Chem, 4(5):709-15 (1996) (alkylated dipeptides).

Identification of modulators through use of the various libraries described herein permits modification of the candidate "hit" (or "lead") to optimize the capacity of the "hit" to bind a polypeptide of the invention. The molecules identified in the binding assay are then tested for antagonist or agonist activity in *in vivo* tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

The binding molecules thus identified may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells such as radioisotopes. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for a polypeptide of the invention. Alternatively, the binding molecules may be complexed with imaging agents for targeting and imaging purposes.

4.10.14 ASSAY FOR RECEPTOR ACTIVITY

The invention also provides methods to detect specific binding of a polypeptide e.g. a ligand or a receptor. The art provides numerous assays particularly useful for identifying previously unknown binding partners for receptor polypeptides of the invention. For example, expression cloning using mammalian or bacterial cells, or dihybrid screening assays can be used to identify polynucleotides encoding binding partners. As another example, affinity chromatography with the appropriate immobilized polypeptide of the invention can be used to isolate polypeptides that recognize and bind polypeptides of the invention. There are a number of different libraries used for the identification of compounds, and in particular small molecules, that modulate (i.e., increase or decrease) biological activity of a polypeptide of the invention. Ligands for receptor polypeptides of the invention can also be identified by adding exogenous ligands, or cocktails of ligands to two cells populations that are genetically identical except for the expression of the receptor of the invention: one cell population expresses the receptor of the invention whereas the other does not. The response of the two cell populations to the addition of ligands(s) are then compared. Alternatively, an expression library can be co-expressed with the polypeptide of the invention in cells and assayed for an autocrine response to identify potential ligand(s). As still another example, BIAcore assays, gel overlay assays, or other methods known in the art can be used to identify binding partner polypeptides, including, (1) organic and inorganic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of random peptides, oligonucleotides or organic molecules.

The role of downstream intracellular signaling molecules in the signaling cascade of the polypeptide of the invention can be determined. For example, a chimeric protein in which the cytoplasmic domain of the polypeptide of the invention is fused to the extracellular portion of a protein, whose ligand has been identified, is produced in a host cell. The cell is then incubated with the ligand specific for the extracellular portion of the chimeric protein, thereby activating the chimeric receptor. Known downstream proteins involved in intracellular signaling can then be assayed for expected modifications i.e. phosphorylation. Other methods known to those in the art can also be used to identify signaling molecules involved in receptor activity.

4.10.15 ANTI-INFLAMMATORY ACTIVITY

Compositions of the present invention may also exhibit anti-inflammatory activity. The anti-inflammatory activity may be achieved by providing a stimulus to cells involved in the inflammatory response, by inhibiting or promoting cell-cell interactions (such as, for example, cell adhesion), by inhibiting or promoting chemotaxis of cells involved in the inflammatory process, inhibiting or promoting cell extravasation, or by stimulating or suppressing production of other factors which more directly inhibit or promote an inflammatory response. Compositions with such activities can be used to treat inflammatory conditions including chronic or acute conditions), including without limitation intimation associated with infection (such as septic shock, sepsis or systemic inflammatory response syndrome (SIRS)), ischemia-reperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine-induced lung injury, inflammatory bowel disease, Crohn's disease or resulting from over production of cytokines such as TNF or IL-1. Compositions of the invention may also be useful to treat anaphylaxis and hypersensitivity to an antigenic substance or material. Compositions of this invention may be utilized to prevent or treat conditions such as, but not limited to, sepsis, acute pancreatitis, endotoxin shock, cytokine induced shock, rheumatoid arthritis, chronic inflammatory arthritis, pancreatic cell damage from diabetes mellitus type 1, graft versus host disease, inflammatory bowel disease, inflamation associated with pulmonary disease, other autoimmune disease or inflammatory disease, an antiproliferative agent such as for acute or chronic mylegenous leukemia or in the prevention of premature labor secondary to intrauterine infections.

4.10.16 LEUKEMIAS

Leukemias and related disorders may be treated or prevented by administration of a therapeutic that promotes or inhibits function of the polynucleotides and/or polypeptides of the invention. Such leukemias and related disorders include but are not limited to acute leukemia,

acute lymphocytic leukemia, acute myelocytic leukemia, myeloblastic, promyelocytic, myelomonocytic, monocytic, erythroleukemia, chronic leukemia, chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia (for a review of such disorders, see Fishman et al., 1985, Medicine, 2d Ed., J.B. Lippincott Co., Philadelphia).

4.10.17 NERVOUS SYSTEM DISORDERS

Nervous system disorders, involving cell types which can be tested for efficacy of intervention with compounds that modulate the activity of the polynucleotides and/or polypeptides of the invention, and which can be treated upon thus observing an indication of therapeutic utility, include but are not limited to nervous system injuries, and diseases or disorders which result in either a disconnection of axons, a diminution or degeneration of neurons, or demyelination. Nervous system lesions which may be treated in a patient (including human and non-human mammalian patients) according to the invention include but are not limited to the following lesions of either the central (including spinal cord, brain) or peripheral nervous systems:

- (i) traumatic lesions, including lesions caused by physical injury or associated with surgery, for example, lesions which sever a portion of the nervous system, or compression injuries;
- (ii) ischemic lesions, in which a lack of oxygen in a portion of the nervous system results in neuronal injury or death, including cerebral infarction or ischemia, or spinal cord infarction or ischemia;
- (iii) infectious lesions, in which a portion of the nervous system is destroyed or injured as a result of infection, for example, by an abscess or associated with infection by human immunodeficiency virus, herpes zoster, or herpes simplex virus or with Lyme disease, tuberculosis, syphilis;
- (iv) degenerative lesions, in which a portion of the nervous system is destroyed or injured as a result of a degenerative process including but not limited to degeneration associated with Parkinson's disease, Alzheimer's disease, Huntington's chorea, or amyotrophic lateral sclerosis;
- (v) lesions associated with nutritional diseases or disorders, in which a portion of the nervous system is destroyed or injured by a nutritional disorder or disorder of metabolism including but not limited to, vitamin B12 deficiency, folic acid deficiency, Wernicke disease, tobacco-alcohol amblyopia, Marchiafava-Bignami disease (primary degeneration of the corpus callosum), and alcoholic cerebellar degeneration;

- (vi) neurological lesions associated with systemic diseases including but not limited to diabetes (diabetic neuropathy, Bell's palsy), systemic lupus erythematosus, carcinoma, or sarcoidosis;
- (vii) lesions caused by toxic substances including alcohol, lead, or particular neurotoxins; and
- (viii) demyelinated lesions in which a portion of the nervous system is destroyed or injured by a demyelinating disease including but not limited to multiple sclerosis, human immunodeficiency virus-associated myelopathy, transverse myelopathy or various etiologies, progressive multifocal leukoencephalopathy, and central pontine myelinolysis.

Therapeutics which are useful according to the invention for treatment of a nervous system disorder may be selected by testing for biological activity in promoting the survival or differentiation of neurons. For example, and not by way of limitation, therapeutics which elicit any of the following effects may be useful according to the invention:

- (i) increased survival time of neurons in culture;
- (ii) increased sprouting of neurons in culture or in vivo;
- (iii) increased production of a neuron-associated molecule in culture or *in vivo*, *e.g.*, choline acetyltransferase or acetylcholinesterase with respect to motor neurons; or
 - (iv) decreased symptoms of neuron dysfunction in vivo.

Such effects may be measured by any method known in the art. In preferred, non-limiting embodiments, increased survival of neurons may be measured by the method set forth in Arakawa et al. (1990, J. Neurosci. 10:3507-3515); increased sprouting of neurons may be detected by methods set forth in Pestronk et al. (1980, Exp. Neurol. 70:65-82) or Brown et al. (1981, Ann. Rev. Neurosci. 4:17-42); increased production of neuron-associated molecules may be measured by bioassay, enzymatic assay, antibody binding, Northern blot assay, etc., depending on the molecule to be measured; and motor neuron dysfunction may be measured by assessing the physical manifestation of motor neuron disorder, e.g., weakness, motor neuron conduction velocity, or functional disability.

In specific embodiments, motor neuron disorders that may be treated according to the invention include but are not limited to disorders such as infarction, infection, exposure to toxin, trauma, surgical damage, degenerative disease or malignancy that may affect motor neurons as well as other components of the nervous system, as well as disorders that selectively affect neurons such as amyotrophic lateral sclerosis, and including but not limited to progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, infantile and juvenile muscular atrophy, progressive bulbar paralysis of childhood (Fazio-Londe syndrome),

poliomyelitis and the post polio syndrome, and Hereditary Motorsensory Neuropathy (Charcot-Marie-Tooth Disease).

4.10.18 OTHER ACTIVITIES

A polypeptide of the invention may also exhibit one or more of the following additional activities or effects: inhibiting the growth, infection or function of, or killing, infectious agents, including, without limitation, bacteria, viruses, fungi and other parasites; effecting (suppressing or enhancing) bodily characteristics, including, without limitation, height, weight, hair color, eye color, skin, fat to lean ratio or other tissue pigmentation, or organ or body part size or shape (such as, for example, breast augmentation or diminution, change in bone form or shape); effecting biorhythms or circadian cycles or rhythms; effecting the fertility of male or female subjects; effecting the metabolism, catabolism, anabolism, processing, utilization, storage or elimination of dietary fat, lipid, protein, carbohydrate, vitamins, minerals, co-factors or other nutritional factors or component(s); effecting behavioral characteristics, including, without limitation, appetite, libido, stress, cognition (including cognitive disorders), depression (including depressive disorders) and violent behaviors; providing analgesic effects or other pain reducing effects; promoting differentiation and growth of embryonic stem cells in lineages other than hematopoietic lineages; hormonal or endocrine activity; in the case of enzymes, correcting deficiencies of the enzyme and treating deficiency-related diseases; treatment of hyperproliferative disorders (such as, for example, psoriasis); immunoglobulin-like activity (such as, for example, the ability to bind antigens or complement); and the ability to act as an antigen in a vaccine composition to raise an immune response against such protein or another material or entity which is cross-reactive with such protein.

4.10.19 IDENTIFICATION OF POLYMORPHISMS

The demonstration of polymorphisms makes possible the identification of such polymorphisms in human subjects and the pharmacogenetic use of this information for diagnosis and treatment. Such polymorphisms may be associated with, e.g., differential predisposition or susceptibility to various disease states (such as disorders involving inflammation or immune response) or a differential response to drug administration, and this genetic information can be used to tailor preventive or therapeutic treatment appropriately. For example, the existence of a polymorphism associated with a predisposition to inflammation or autoimmune disease makes possible the diagnosis of this condition in humans by identifying the presence of the polymorphism.

Polymorphisms can be identified in a variety of ways known in the art which all generally involve obtaining a sample from a patient, analyzing DNA from the sample, optionally involving isolation or amplification of the DNA, and identifying the presence of the polymorphism in the DNA. For example, PCR may be used to amplify an appropriate fragment of genomic DNA which may then be sequenced. Alternatively, the DNA may be subjected to allele-specific oligonucleotide hybridization (in which appropriate oligonucleotides are hybridized to the DNA under conditions permitting detection of a single base mismatch) or to a single nucleotide extension assay (in which an oligonucleotide that hybridizes immediately adjacent to the position of the polymorphism is extended with one or more labeled nucleotides). In addition, traditional restriction fragment length polymorphism analysis (using restriction enzymes that provide differential digestion of the genomic DNA depending on the presence or absence of the polymorphism) may be performed. Arrays with nucleotide sequences of the present invention can be used to detect polymorphisms. The array can comprise modified nucleotide sequences of the present invention in order to detect the nucleotide sequences of the present invention. In the alternative, any one of the nucleotide sequences of the present invention can be placed on the array to detect changes from those sequences.

Alternatively a polymorphism resulting in a change in the amino acid sequence could also be detected by detecting a corresponding change in amino acid sequence of the protein, e.g., by an antibody specific to the variant sequence.

4.10.20 ARTHRITIS AND INFLAMMATION

The immunosuppressive effects of the compositions of the invention against rheumatoid arthritis is determined in an experimental animal model system. The experimental model system is adjuvant induced arthritis in rats, and the protocol is described by J. Holoshitz, et at., 1983, Science, 219:56, or by B. Waksman et al., 1963, Int. Arch. Allergy Appl. Immunol., 23:129. Induction of the disease can be caused by a single injection, generally intradermally, of a suspension of killed Mycobacterium tuberculosis in complete Freund's adjuvant (CFA). The route of injection can vary, but rats may be injected at the base of the tail with an adjuvant mixture. The polypeptide is administered in phosphate buffered solution (PBS) at a dose of about 1-5 mg/kg. The control consists of administering PBS only.

The procedure for testing the effects of the test compound would consist of intradermally injecting killed Mycobacterium tuberculosis in CFA followed by immediately administering the test compound and subsequent treatment every other day until day 24. At 14, 15, 18, 20, 22, and 24 days after injection of Mycobacterium CFA, an overall arthritis score may be obtained as described by J. Holoskitz above. An analysis of the data would reveal that the test compound

would have a dramatic affect on the swelling of the joints as measured by a decrease of the arthritis score.

4.11 THERAPEUTIC METHODS

The compositions (including polypeptide fragments, analogs, variants and antibodies or other binding partners or modulators including antisense polynucleotides) of the invention have numerous applications in a variety of therapeutic methods. Examples of therapeutic applications include, but are not limited to, those exemplified herein.

4.11.1 EXAMPLE

One embodiment of the invention is the administration of an effective amount of the polypeptides or other composition of the invention to individuals affected by a disease or disorder that can be modulated by regulating the peptides of the invention. While the mode of administration is not particularly important, parenteral administration is preferred. An exemplary mode of administration is to deliver an intravenous bolus. The dosage of the polypeptides or other composition of the invention will normally be determined by the prescribing physician. It is to be expected that the dosage will vary according to the age, weight, condition and response of the individual patient. Typically, the amount of polypeptide administered per dose will be in the range of about 0.01µg/kg to 100 mg/kg of body weight, with the preferred dose being about 0.1µg/kg to 10 mg/kg of patient body weight. For parenteral administration, polypeptides of the invention will be formulated in an injectable form combined with a pharmaceutically acceptable parenteral vehicle. Such vehicles are well known in the art and examples include water, saline, Ringer's solution, dextrose solution, and solutions consisting of small amounts of the human serum albumin. The vehicle may contain minor amounts of additives that maintain the isotonicity and stability of the polypeptide or other active ingredient. The preparation of such solutions is within the skill of the art.

4.12 PHARMACEUTICAL FORMULATIONS AND ROUTES OF ADMINISTRATION

A protein or other composition of the present invention (from whatever source derived, including without limitation from recombinant and non-recombinant sources and including antibodies and other binding partners of the polypeptides of the invention) may be administered to a patient in need, by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s) at doses to treat or ameliorate a variety of disorders. Such a composition may optionally contain (in addition to protein or other active ingredient and a carrier) diluents,

fillers, salts, buffers, stabilizers, solubilizers, and other materials well known in the art. The term "pharmaceutically acceptable" means a non-toxic material that does not interfere with the effectiveness of the biological activity of the active ingredient(s). The characteristics of the carrier will depend on the route of administration. The pharmaceutical composition of the invention may also contain cytokines, lymphokines, or other hematopoietic factors such as M-CSF, GM-CSF, TNF, IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12. IL-13, IL-14, IL-15, IFN, TNF0, TNF1, TNF2, G-CSF, Meg-CSF, thrombopoietin, stem cell factor, and erythropoietin. In further compositions, proteins of the invention may be combined with other agents beneficial to the treatment of the disease or disorder in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet-derived growth factor (PDGF), transforming growth factors (TGF-α and TGF-β), insulin-like growth factor (IGF), as well as cytokines described herein.

The pharmaceutical composition may further contain other agents which either enhance the activity of the protein or other active ingredient or complement its activity or use in treatment. Such additional factors and/or agents may be included in the pharmaceutical composition to produce a synergistic effect with protein or other active ingredient of the invention, or to minimize side effects. Conversely, protein or other active ingredient of the present invention may be included in formulations of the particular clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent to minimize side effects of the clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent (such as IL-1Ra, IL-1 Hy1, IL-1 Hy2, anti-TNF, corticosteroids, immunosuppressive agents). A protein of the present invention may be active in multimers (e.g., heterodimers or homodimers) or complexes with itself or other proteins. As a result, pharmaceutical compositions of the invention may comprise a protein of the invention in such multimeric or complexed form.

As an alternative to being included in a pharmaceutical composition of the invention including a first protein, a second protein or a therapeutic agent may be concurrently administered with the first protein (e.g., at the same time, or at differing times provided that therapeutic concentrations of the combination of agents is achieved at the treatment site). Techniques for formulation and administration of the compounds of the instant application may be found in "Remington's Pharmaceutical Sciences," Mack Publishing Co., Easton, PA, latest edition. A therapeutically effective dose further refers to that amount of the compound sufficient to result in amelioration of symptoms, e.g., treatment, healing, prevention or amelioration of the relevant medical condition, or an increase in rate of treatment, healing, prevention or amelioration of such conditions. When applied to an individual active ingredient, administered

alone, a therapeutically effective dose refers to that ingredient alone. When applied to a combination, a therapeutically effective dose refers to combined amounts of the active ingredients that result in the therapeutic effect, whether administered in combination, serially or simultaneously.

In practicing the method of treatment or use of the present invention, a therapeutically effective amount of protein or other active ingredient of the present invention is administered to a mammal having a condition to be treated. Protein or other active ingredient of the present invention may be administered in accordance with the method of the invention either alone or in combination with other therapies such as treatments employing cytokines, lymphokines or other hematopoietic factors. When co- administered with one or more cytokines, lymphokines or other hematopoietic factors, protein or other active ingredient of the present invention may be administered either simultaneously with the cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors, or sequentially. If administered sequentially, the attending physician will decide on the appropriate sequence of administering protein or other active ingredient of the present invention in combination with cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors.

4.12.1 ROUTES OF ADMINISTRATION

Suitable routes of administration may, for example, include oral, rectal, transmucosal, or intestinal administration; parenteral delivery, including intramuscular, subcutaneous, intramedullary injections, as well as intrathecal, direct intraventricular, intravenous, intraperitoneal, intranasal, or intraocular injections. Administration of protein or other active ingredient of the present invention used in the pharmaceutical composition or to practice the method of the present invention can be carried out in a variety of conventional ways, such as oral ingestion, inhalation, topical application or cutaneous, subcutaneous, intraperitoneal, parenteral or intravenous injection. Intravenous administration to the patient is preferred.

Alternately, one may administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into a arthritic joints or in fibrotic tissue, often in a depot or sustained release formulation. In order to prevent the scarring process frequently occurring as complication of glaucoma surgery, the compounds may be administered topically, for example, as eye drops. Furthermore, one may administer the drug in a targeted drug delivery system, for example, in a liposome coated with a specific antibody, targeting, for example, arthritic or fibrotic tissue. The liposomes will be targeted to and taken up selectively by the afflicted tissue.

The polypeptides of the invention are administered by any route that delivers an effective dosage to the desired site of action. The determination of a suitable route of administration and an effective dosage for a particular indication is within the level of skill in the art. Preferably for wound treatment, one administers the therapeutic compound directly to the site. Suitable dosage ranges for the polypeptides of the invention can be extrapolated from these dosages or from similar studies in appropriate animal models. Dosages can then be adjusted as necessary by the clinician to provide maximal therapeutic benefit.

4.12.2 COMPOSITIONS/FORMULATIONS

Pharmaceutical compositions for use in accordance with the present invention thus may be formulated in a conventional manner using one or more physiologically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. These pharmaceutical compositions may be manufactured in a manner that is itself known, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Proper formulation is dependent upon the route of administration chosen. When a therapeutically effective amount of protein or other active ingredient of the present invention is administered orally, protein or other active ingredient of the present Invention will be in the form of a tablet, capsule, powder, solution or elixir. When administered in tablet form, the pharmaceutical composition of the invention may additionally contain a solid carrier such as a gelatin or an adjuvant. The tablet, capsule, and powder contain from about 5 to 95% protein or other active ingredient of the present invention, and preferably from about 25 to 90% protein or other active ingredient of the present invention. When administered in liquid form, a liquid carrier such as water, petroleum, oils of animal or plant origin such as peanut oil, mineral oil, soybean oil, or sesame oil, or synthetic oils may be added. The liquid form of the pharmaceutical composition may further contain physiological saline solution, dextrose or other saccharide solution, or glycols such as ethylene glycol, propylene glycol or polyethylene glycol. When administered in liquid form, the pharmaceutical composition contains from about 0.5 to 90% by weight of protein or other active ingredient of the present invention, and preferably from about 1 to 50% protein or other active ingredient of the present invention.

When a therapeutically effective amount of protein or other active ingredient of the present invention is administered by intravenous, cutaneous or subcutaneous injection, protein or other active ingredient of the present invention will be in the form of a pyrogen-free, parenterally acceptable aqueous solution. The preparation of such parenterally acceptable protein or other active ingredient solutions, having due regard to pH, isotonicity, stability, and the like, is within

the skill in the art. A preferred pharmaceutical composition for intravenous, cutaneous, or subcutaneous injection should contain, in addition to protein or other active ingredient of the present invention, an isotonic vehicle such as Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, Lactated Ringer's Injection, or other vehicle as known in the art. The pharmaceutical composition of the present invention may also contain stabilizers, preservatives, buffers, antioxidants, or other additives known to those of skill in the art. For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks's solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

For oral administration, the compounds can be formulated readily by combining the active compounds with pharmaceutically acceptable carriers well known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated. Pharmaceutical preparations for oral use can be obtained from a solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate. Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as tale or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition,

stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration. For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebuliser, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g., gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch. The compounds may be formulated for parenteral administration by injection, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active compounds in water-soluble form. Additionally, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

The compounds may also be formulated in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glycerides. In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

A pharmaceutical carrier for the hydrophobic compounds of the invention is a co-solvent system comprising benzyl alcohol, a nonpolar surfactant, a water-miscible organic polymer, and an aqueous phase. The co-solvent system may be the VPD co-solvent system. VPD is a solution of 3% w/v benzyl alcohol, 8% w/v of the nonpolar surfactant polysorbate 80, and 65% w/v polyethylene glycol 300, made up to volume in absolute ethanol. The VPD co-solvent system (VPD:5W) consists of VPD diluted 1:1 with a 5% dextrose in water solution. This co-solvent system dissolves hydrophobic compounds well, and itself produces low toxicity upon systemic administration. Naturally, the proportions of a co-solvent system may be varied considerably without destroying its solubility and toxicity characteristics. Furthermore, the identity of the co-solvent components may be varied: for example, other low-toxicity nonpolar surfactants may be used instead of polysorbate 80; the fraction size of polyethylene glycol may be varied; other biocompatible polymers may replace polyethylene glycol, e.g. polyvinyl pyrrolidone; and other sugars or polysaccharides may substitute for dextrose. Alternatively, other delivery systems for hydrophobic pharmaceutical compounds may be employed. Liposomes and emulsions are well known examples of delivery vehicles or carriers for hydrophobic drugs. Certain organic solvents such as dimethylsulfoxide also may be employed, although usually at the cost of greater toxicity. Additionally, the compounds may be delivered using a sustained-release system, such as semipermeable matrices of solid hydrophobic polymers containing the therapeutic agent. Various types of sustained-release materials have been established and are well known by those skilled in the art. Sustained-release capsules may, depending on their chemical nature, release the compounds for a few weeks up to over 100 days. Depending on the chemical nature and the biological stability of the therapeutic reagent, additional strategies for protein or other active ingredient stabilization may be employed.

The pharmaceutical compositions also may comprise suitable solid or gcl phase carriers or excipients. Examples of such carriers or excipients include but are not limited to calcium carbonate, calcium phosphate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols. Many of the active ingredients of the invention may be provided as salts with pharmaceutically compatible counter ions. Such pharmaceutically acceptable base addition salts are those salts which retain the biological effectiveness and properties of the free acids and which are obtained by reaction with inorganic or organic bases such as sodium hydroxide, magnesium hydroxide, ammonia, trialkylamine, dialkylamine, monoalkylamine, dibasic amino acids, sodium acetate, potassium benzoate, triethanol amine and the like.

The pharmaceutical composition of the invention may be in the form of a complex of the protein(s) or other active ingredient(s) of present invention along with protein or peptide

antigens. The protein and/or peptide antigen will deliver a stimulatory signal to both B and T lymphocytes. B lymphocytes will respond to antigen through their surface immunoglobulin receptor. T lymphocytes will respond to antigen through the T cell receptor (TCR) following presentation of the antigen by MHC proteins. MHC and structurally related proteins including those encoded by class I and class II MHC genes on host cells will serve to present the peptide antigen(s) to T lymphocytes. The antigen components could also be supplied as purified MHC-peptide complexes alone or with co-stimulatory molecules that can directly signal T cells. Alternatively antibodies able to bind surface immunoglobulin and other molecules on B cells as well as antibodies able to bind the TCR and other molecules on T cells can be combined with the pharmaceutical composition of the invention.

The pharmaceutical composition of the invention may be in the form of a liposome in which protein of the present invention is combined, in addition to other pharmaceutically acceptable carriers, with amphipathic agents such as lipids which exist in aggregated form as micelles, insoluble monolayers, liquid crystals, or lamellar layers in aqueous solution. Suitable lipids for liposomal formulation include, without limitation, monoglycerides, diglycerides, sulfatides, lysolecithins, phospholipids, saponin, bile acids, and the like. Preparation of such liposomal formulations is within the level of skill in the art, as disclosed, for example, in U.S. Patent Nos. 4,235,871; 4,501,728; 4,837,028; and 4,737,323, all of which are incorporated herein by reference.

The amount of protein or other active ingredient of the present invention in the pharmaceutical composition of the present invention will depend upon the nature and severity of the condition being treated, and on the nature of prior treatments which the patient has undergone. Ultimately, the attending physician will decide the amount of protein or other active ingredient of the present invention with which to treat each individual patient. Initially, the attending physician will administer low doses of protein or other active ingredient of the present invention and observe the patient's response. Larger doses of protein or other active ingredient of the present invention may be administered until the optimal therapeutic effect is obtained for the patient, and at that point the dosage is not increased further. It is contemplated that the various pharmaceutical compositions used to practice the method of the present invention should contain about 0.01 µg to about 100 mg (preferably about 0.1 µg to about 10 mg, more preferably about 0.1 µg to about 1 mg) of protein or other active ingredient of the present invention per kg body weight. For compositions of the present invention which are useful for bone, cartilage, tendon or ligament regeneration, the therapeutic method includes administering the composition topically, systematically, or locally as an implant or device. When administered, the therapeutic composition for use in this invention is, of course, in a pyrogen-free, physiologically acceptable

form. Further, the composition may desirably be encapsulated or injected in a viscous form for delivery to the site of bone, cartilage or tissue damage. Topical administration may be suitable for wound healing and tissue repair. Therapeutically useful agents other than a protein or other active ingredient of the invention which may also optionally be included in the composition as described above, may alternatively or additionally, be administered simultaneously or sequentially with the composition in the methods of the invention. Preferably for bone and/or cartilage formation, the composition would include a matrix capable of delivering the protein-containing or other active ingredient-containing composition to the site of bone and/or cartilage damage, providing a structure for the developing bone and cartilage and optimally capable of being resorbed into the body. Such matrices may be formed of materials presently in use for other implanted medical applications.

The choice of matrix material is based on biocompatibility, biodegradability, mechanical properties, cosmetic appearance and interface properties. The particular application of the compositions will define the appropriate formulation. Potential matrices for the compositions may be biodegradable and chemically defined calcium sulfate, tricalcium phosphate, hydroxyapatite, polylactic acid, polyglycolic acid and polyanhydrides. Other potential materials are biodegradable and biologically well-defined, such as bone or dermal collagen. Further matrices are comprised of pure proteins or extracellular matrix components. Other potential matrices are nonbiodegradable and chemically defined, such as sintered hydroxyapatite, bioglass, aluminates, or other ceramics. Matrices may be comprised of combinations of any of the above mentioned types of material, such as polylactic acid and hydroxyapatite or collagen and tricalcium phosphate. The bioceramics may be altered in composition, such as in calcium-aluminate-phosphate and processing to alter pore size, particle size, particle shape, and biodegradability. Presently preferred is a 50:50 (mole weight) copolymer of lactic acid and glycolic acid in the form of porous particles having diameters ranging from 150 to 800 microns. In some applications, it will be useful to utilize a sequestering agent, such as carboxymethyl cellulose or autologous blood clot, to prevent the protein compositions from disassociating from the matrix.

A preferred family of sequestering agents is cellulosic materials such as alkylcelluloses (including hydroxyalkylcelluloses), including methylcellulose, ethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropyl-methylcellulose, and carboxymethylcellulose, the most preferred being cationic salts of carboxymethylcellulose (CMC). Other preferred sequestering agents include hyaluronic acid, sodium alginate, poly(ethylene glycol), polyoxyethylene oxide, carboxyvinyl polymer and poly(vinyl alcohol). The amount of sequestering agent useful herein is 0.5-20 wt %, preferably 1-10 wt % based on

total formulation weight, which represents the amount necessary to prevent desorption of the protein from the polymer matrix and to provide appropriate handling of the composition, yet not so much that the progenitor cells are prevented from infiltrating the matrix, thereby providing the protein the opportunity to assist the osteogenic activity of the progenitor cells. In further compositions, proteins or other active ingredients of the invention may be combined with other agents beneficial to the treatment of the bone and/or cartilage defect, wound, or tissue in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet derived growth factor (PDGF), transforming growth factors (TGF- α and TGF- β), and insulin-like growth factor (IGF).

The therapeutic compositions are also presently valuable for veterinary applications. Particularly domestic animals and thoroughbred horses, in addition to humans, are desired patients for such treatment with proteins or other active ingredients of the present invention. The dosage regimen of a protein-containing pharmaceutical composition to be used in tissue regeneration will be determined by the attending physician considering various factors which modify the action of the proteins, e.g., amount of tissue weight desired to be formed, the site of damage, the condition of the damaged tissue, the size of a wound, type of damaged tissue (e.g., bone), the patient's age, sex, and diet, the severity of any infection, time of administration and other clinical factors. The dosage may vary with the type of matrix used in the reconstitution and with inclusion of other proteins in the pharmaceutical composition. For example, the addition of other known growth factors, such as IGF I (insulin like growth factor I), to the final composition, may also effect the dosage. Progress can be monitored by periodic assessment of tissue/bone growth and/or repair, for example, X-rays, histomorphometric determinations and tetracycline labeling.

Polynucleotides of the present invention can also be used for gene therapy. Such polynucleotides can be introduced either in vivo or ex vivo into cells for expression in a mammalian subject. Polynucleotides of the invention may also be administered by other known methods for introduction of nucleic acid into a cell or organism (including, without limitation, in the form of viral vectors or naked DNA). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes.

4.12.3 EFFECTIVE DOSAGE

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve its intended purpose. More specifically, a therapeutically effective amount means an amount

effective to prevent development of or to alleviate the existing symptoms of the subject being treated. Determination of the effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein. For any compound used in the method of the invention, the therapeutically effective dose can be estimated initially from appropriate in vitro assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that can be used to more accurately determine useful doses in humans. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes the IC₅₀ as determined in cell culture (*i.e.*, the concentration of the test compound which achieves a half-maximal inhibition of the protein's biological activity). Such information can be used to more accurately determine useful doses in humans.

A therapeutically effective dose refers to that amount of the compound that results in amelioration of symptoms or a prolongation of survival in a patient. Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD50 (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio between LD₅₀ and ED₅₀. Compounds which exhibit high therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in human. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED50 with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. See, e.g., Fingl et al., 1975, in "The Pharmacological Basis of Therapeutics", Ch. 1 p.1. Dosage amount and interval may be adjusted individually to provide plasma levels of the active moiety which are sufficient to maintain the desired effects, or minimal effective concentration (MEC). The MEC will vary for each compound but can be estimated from in vitro data. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. However, HPLC assays or bioassays can be used to determine plasma concentrations.

Dosage intervals can also be determined using MEC value. Compounds should be administered using a regimen which maintains plasma levels above the MEC for 10-90% of the time, preferably between 30-90% and most preferably between 50-90%. In cases of local administration or selective uptake, the effective local concentration of the drug may not be related to plasma concentration.

An exemplary dosage regimen for polypeptides or other compositions of the invention will be in the range of about $0.01~\mu g/kg$ to 100~mg/kg of body weight daily, with the preferred dose being about $0.1~\mu g/kg$ to 25~mg/kg of patient body weight daily, varying in adults and children. Dosing may be once daily, or equivalent doses may be delivered at longer or shorter intervals.

The amount of composition administered will, of course, be dependent on the subject being treated, on the subject's age and weight, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.

4.12.4 PACKAGING

The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The pack may, for example, comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration. Compositions comprising a compound of the invention formulated in a compatible pharmaceutical carrier may also be prepared, placed in an appropriate container, and labeled for treatment of an indicated condition.

4.13 ANTIBODIES

Also included in the invention are antibodies to proteins, or fragments of proteins of the invention. The term "antibody" as used herein refers to immunoglobulin molecules and immunologically active portions of immunoglobulin (Ig) molecules, i.e., molecules that contain an antigen binding site that specifically binds (immunoreacts with) an antigen. Such antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, F_{ab} , $F_{ab'}$ and $F_{(ab')2}$ fragments, and an F_{ab} expression library. In general, an antibody molecule obtained from humans relates to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes have subclasses as well, such as IgG₁, IgG₂, and others. Furthermore, in humans, the light chain may be a kappa chain or a lambda chain. Reference herein to antibodies includes a reference to all such classes, subclasses and types of human antibody species.

An isolated related protein of the invention may be intended to serve as an antigen, or a portion or fragment thereof, and additionally can be used as an immunogen to generate antibodies that immunospecifically bind the antigen, using standard techniques for polyclonal and monoclonal antibody preparation. The full-length protein can be used or, alternatively, the invention provides antigenic peptide fragments of the antigen for use as immunogens. An antigenic peptide fragment comprises at least 6 amino acid residues of the amino acid sequence

of the full length protein, such as the amino acid sequences shown in SEQ ID NO: 445-888, and encompasses an epitope thereof such that an antibody raised against the peptide forms a specific immune complex with the full length protein or with any fragment that contains the epitope. Preferably, the antigenic peptide comprises at least 10 amino acid residues, or at least 15 amino acid residues, or at least 20 amino acid residues. Preferred epitopes encompassed by the antigenic peptide are regions of the protein that are located on its surface; commonly these are hydrophilic regions.

In certain embodiments of the invention, at least one epitope encompassed by the antigenic peptide is a region of -related protein that is located on the surface of the protein, e.g., a hydrophilic region. A hydrophobicity analysis of the human related protein sequence will indicate which regions of a related protein are particularly hydrophilic and, therefore, are likely to encode surface residues useful for targeting antibody production. As a means for targeting antibody production, hydropathy plots showing regions of hydrophilicity and hydrophobicity may be generated by any method well known in the art, including, for example, the Kyte Doolittle or the Hopp Woods methods, either with or without Fourier transformation. See, e.g., Hopp and Woods, 1981, Proc. Nat. Acad. Sci. USA 78: 3824-3828; Kyte and Doolittle 1982, J. Mol. Biol. 157: 105-142, each of which is incorporated herein by reference in its entirety. Antibodies that are specific for one or more domains within an antigenic protein, or derivatives, fragments, analogs or homologs thereof, are also provided herein.

A protein of the invention, or a derivative, fragment, analog, homolog or ortholog thereof, may be utilized as an immunogen in the generation of antibodies that immunospecifically bind these protein components.

Various procedures known within the art may be used for the production of polyclonal or monoclonal antibodies directed against a protein of the invention, or against derivatives, fragments, analogs homologs or orthologs thereof (see, for example, Antibodies: A Laboratory Manual, Harlow E, and Lane D, 1988, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, incorporated herein by reference). Some of these antibodies are discussed below.

4.13.1 POLYCLONAL ANTIBODIES

For the production of polyclonal antibodies, various suitable host animals (e.g., rabbit, goat, mouse or other mammal) may be immunized by one or more injections with the native protein, a synthetic variant thereof, or a derivative of the foregoing. An appropriate immunogenic preparation can contain, for example, the naturally occurring immunogenic protein, a chemically synthesized polypeptide representing the immunogenic protein, or a recombinantly expressed immunogenic protein. Furthermore, the protein may be conjugated to

a second protein known to be immunogenic in the mammal being immunized. Examples of such immunogenic proteins include but are not limited to keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. The preparation can further include an adjuvant. Various adjuvants used to increase the immunological response include, but are not limited to, Freund's (complete and incomplete), mineral gels (e.g., aluminum hydroxide), surface active substances (e.g., lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, dinitrophenol, etc.), adjuvants usable in humans such as Bacille Calmette-Guerin and Corynebacterium parvum, or similar immunostimulatory agents. Additional examples of adjuvants which can be employed include MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate).

The polyclonal antibody molecules directed against the immunogenic protein can be isolated from the mammal (e.g., from the blood) and further purified by well known techniques, such as affinity chromatography using protein A or protein G, which provide primarily the IgG fraction of immune serum. Subsequently, or alternatively, the specific antigen which is the target of the immunoglobulin sought, or an epitope thereof, may be immobilized on a column to purify the immune specific antibody by immunoaffinity chromatography. Purification of immunoglobulins is discussed, for example, by D. Wilkinson (The Scientist, published by The Scientist, Inc., Philadelphia PA, Vol. 14, No. 8 (April 17, 2000), pp. 25-28).

4.13.2 MONOCLONAL ANTIBODIES

The term "monoclonal antibody" (MAb) or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs thus contain an antigen binding site capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it.

Monoclonal antibodies can be prepared using hybridoma methods, such as those described by Kohler and Milstein, Nature, 256:495 (1975). In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes can be immunized in vitro. The immunizing agent will typically include the protein antigen, a fragment thereof or a fusion protein thereof. Generally, either peripheral blood lymphocytes are used if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are

desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding, Monoclonal Antibodies: Principles and Practice, Academic Press, (1986) pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovine and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells can be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing cells, and are sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture Collection, Manassas, Virginia. Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, J. Immunol., 133:3001 (1984); Brodeur et al., Monoclonal Antibody Production Techniques and Applications, Marcel Dekker, Inc., New York, (1987) pp. 51-63).

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against the antigen. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, Anal. Biochem., 107:220 (1980). Preferably, antibodies having a high degree of specificity and a high binding affinity for the target antigen are isolated.

After the desired hybridoma cells are identified, the clones can be subcloned by limiting dilution procedures and grown by standard methods. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium. Alternatively, the hybridoma cells can be grown in vivo as ascites in a mammal. The monoclonal antibodies secreted by the subclones can be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, for

example, protein A-Sepharose, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies can also be made by recombinant DNA methods, such as those described in U.S. Patent No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA can be placed into expression vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also can be modified, for example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences (U.S. Patent No. 4,816,567; Morrison, Nature 368, 812-13 (1994)) or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

4.13.3 HUMANIZED ANTIBODIES

The antibodies directed against the protein antigens of the invention can further comprise humanized antibodies or human antibodies. These antibodies are suitable for administration to humans without engendering an immune response by the human against the administered immunoglobulin. Humanized forms of antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigenbinding subsequences of antibodies) that are principally comprised of the sequence of a human immunoglobulin, and contain minimal sequence derived from a non-human immunoglobulin. Humanization can be performed following the method of Winter and co-workers (Jones et al., Nature, 321:522-525 (1986); Riechmann et al., Nature, 332:323-327 (1988); Verhoeyen et al., Science, 239:1534-1536 (1988)), by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. (See also U.S. Patent No. 5,225,539.) In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies can also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable

domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., 1986; Riechmann et al., 1988; and Presta, Curr. Op. Struct. Biol., 2:593-596 (1992)).

4.13.4 HUMAN ANTIBODIES

Fully human antibodies relate to antibody molecules in which essentially the entire sequences of both the light chain and the heavy chain, including the CDRs, arise from human genes. Such antibodies are termed "human antibodies", or "fully human antibodies" herein. Human monoclonal antibodies can be prepared by the trioma technique; the human B-cell hybridoma technique (see Kozbor, et al., 1983 Immunol Today 4: 72) and the EBV hybridoma technique to produce human monoclonal antibodies (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96). Human monoclonal antibodies may be utilized in the practice of the present invention and may be produced by using human hybridomas (see Cote, et al., 1983. Proc Natl Acad Sci USA 80: 2026-2030) or by transforming human B-cells with Epstein Barr Virus in vitro (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96).

In addition, human antibodies can also be produced using additional techniques, including phage display libraries (Hoogenboom and Winter, J. Mol. Biol., 227:381 (1991); Marks et al., J. Mol. Biol., 222:581 (1991)). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in all respects, including gene rearrangement, assembly, and antibody repertoire. This approach is described, for example, in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,016, and in Marks et al. (Bio/Technology 10, 779-783 (1992)); Lonberg et al. (Nature 368 856-859 (1994)); Morrison (Nature 368, 812-13 (1994)); Fishwild et al. (Nature Biotechnology 14, 845-51 (1996)); Neuberger (Nature Biotechnology 14, 826 (1996)); and Lonberg and Huszar (Intern. Rev. Immunol. 13 65-93 (1995)).

Human antibodies may additionally be produced using transgenic nonhuman animals which are modified so as to produce fully human antibodies rather than the animal's endogenous antibodies in response to challenge by an antigen. (See PCT publication WO94/02602). The endogenous genes encoding the heavy and light immunoglobulin chains in the nonhuman host

have been incapacitated, and active loci encoding human heavy and light chain immunoglobulins are inserted into the host's genome. The human genes are incorporated, for example, using yeast artificial chromosomes containing the requisite human DNA segments. An animal which provides all the desired modifications is then obtained as progeny by crossbreeding intermediate transgenic animals containing fewer than the full complement of the modifications. The preferred embodiment of such a nonhuman animal is a mouse, and is termed the XenomouseTM as disclosed in PCT publications WO 96/33735 and WO 96/34096. This animal produces B cells which secrete fully human immunoglobulins. The antibodies can be obtained directly from the animal after immunization with an immunogen of interest, as, for example, a preparation of a polyclonal antibody, or alternatively from immortalized B cells derived from the animal, such as hybridomas producing monoclonal antibodies. Additionally, the genes encoding the immunoglobulins with human variable regions can be recovered and expressed to obtain the antibodies directly, or can be further modified to obtain analogs of antibodies such as, for example, single chain Fv molecules.

An example of a method of producing a nonhuman host, exemplified as a mouse, lacking expression of an endogenous immunoglobulin heavy chain is disclosed in U.S. Patent No. 5,939,598. It can be obtained by a method including deleting the J segment genes from at least one endogenous heavy chain locus in an embryonic stem cell to prevent rearrangement of the locus and to prevent formation of a transcript of a rearranged immunoglobulin heavy chain locus, the deletion being effected by a targeting vector containing a gene encoding a selectable marker; and producing from the embryonic stem cell a transgenic mouse whose somatic and germ cells contain the gene encoding the selectable marker.

A method for producing an antibody of interest, such as a human antibody, is disclosed in U.S. Patent No. 5,916,771. It includes introducing an expression vector that contains a nucleotide sequence encoding a heavy chain into one mammalian host cell in culture, introducing an expression vector containing a nucleotide sequence encoding a light chain into another mammalian host cell, and fusing the two cells to form a hybrid cell. The hybrid cell expresses an antibody containing the heavy chain and the light chain.

In a further improvement on this procedure, a method for identifying a clinically relevant epitope on an immunogen, and a correlative method for selecting an antibody that binds immunospecifically to the relevant epitope with high affinity, are disclosed in PCT publication WO 99/53049.

4.13.5 Fab FRAGMENTS AND SINGLE CHAIN ANTIBODIES

According to the invention, techniques can be adapted for the production of single-chain antibodies specific to an antigenic protein of the invention (see e.g., U.S. Patent No. 4,946,778). In addition, methods can be adapted for the construction of F_{ab} expression libraries (see e.g., Huse, et al., 1989 Science 246: 1275-1281) to allow rapid and effective identification of monoclonal F_{ab} fragments with the desired specificity for a protein or derivatives, fragments, analogs or homologs thereof. Antibody fragments that contain the idiotypes to a protein antigen may be produced by techniques known in the art including, but not limited to: (i) an $F_{(ab')2}$ fragment produced by pepsin digestion of an antibody molecule; (ii) an F_{ab} fragment generated by reducing the disulfide bridges of an $F_{(ab')2}$ fragment; (iii) an F_{ab} fragment generated by the treatment of the antibody molecule with papain and a reducing agent and (iv) F_{v} fragments.

4.13.6 BISPECIFIC ANTIBODIES

Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens. In the present case, one of the binding specificities is for an antigenic protein of the invention. The second binding target is any other antigen, and advantageously is a cell-surface protein or receptor or receptor subunit.

Methods for making bispecific antibodies are known in the art. Traditionally, the recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have different specificities (Milstein and Cuello, Nature, 305:537-539 (1983)). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traunecker *et al.*, 1991 *EMBO J.*, 10:3655-3659.

Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host organism. For further details of generating bispecific antibodies see, for example, Suresh et al., Methods in Enzymology, 121:210 (1986).

According to another approach described in WO 96/27011, the interface between a pair of antibody molecules can be engineered to maximize the percentage of heterodimers which are recovered from recombinant cell culture. The preferred interface comprises at least a part of the CH3 region of an antibody constant domain. In this method, one or more small amino acid side chains from the interface of the first antibody molecule are replaced with larger side chains (e.g. tyrosine or tryptophan). Compensatory "cavities" of identical or similar size to the large side chain(s) are created on the interface of the second antibody molecule by replacing large amino acid side chains with smaller ones (e.g. alanine or threonine). This provides a mechanism for increasing the yield of the heterodimer over other unwanted end-products such as homodimers.

Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g. F(ab')₂ bispecific antibodies). Techniques for generating bispecific antibodies from antibody fragments have been described in the literature. For example, bispecific antibodies can be prepared using chemical linkage. Brennan et al., Science 229:81 (1985) describe a procedure wherein intact antibodies are proteolytically cleaved to generate F(ab')₂ fragments. These fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The Fab' fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the Fab'-TNB derivatives is then reconverted to the Fab'-thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other Fab'-TNB derivative to form the bispecific antibody. The bispecific antibodies produced can be used as agents for the selective immobilization of enzymes.

Additionally, Fab' fragments can be directly recovered from E. coli and chemically coupled to form bispecific antibodies. Shalaby et al., J. Exp. Med. 175:217-225 (1992) describe the production of a fully humanized bispecific antibody F(ab')₂ molecule. Each Fab' fragment was separately secreted from E. coli and subjected to directed chemical coupling in vitro to form the bispecific antibody. The bispecific antibody thus formed was able to bind to cells overexpressing the ErbB2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets.

Various techniques for making and isolating bispecific antibody fragments directly from recombinant cell culture have also been described. For example, bispecific antibodies have been produced using leucine zippers. Kostelny et al., <u>J. Immunol.</u> 148(5):1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins were linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers were reduced at the hinge region to form monomers and then re-oxidized to form the antibody heterodimers. This method can also be utilized for the production of antibody homodimers. The "diabody" technology

described by Hollinger et al., <u>Proc. Natl. Acad. Sci. USA</u> 90:6444-6448 (1993) has provided an alternative mechanism for making bispecific antibody fragments. The fragments comprise a heavy-chain variable domain (V_H) connected to a light-chain variable domain (V_L) by a linker which is too short to allow pairing between the two domains on the same chain. Accordingly, the V_H and V_L domains of one fragment are forced to pair with the complementary V_L and V_H domains of another fragment, thereby forming two antigen-binding sites. Another strategy for making bispecific antibody fragments by the use of single-chain Fv (sFv) dimers has also been reported. See, Gruber et al., <u>J. Immunol.</u> 152:5368 (1994).

Antibodies with more than two valencies are contemplated. For example, trispecific antibodies can be prepared. Tutt et al., J. Immunol. 147:60 (1991). Exemplary bispecific antibodies can bind to two different epitopes, at least one of which originates in the protein antigen of the invention. Alternatively, an anti-antigenic arm of an immunoglobulin molecule can be combined with an arm which binds to a triggering molecule on a leukocyte such as a T-cell receptor molecule (e.g. CD2, CD3, CD28, or B7), or Fc receptors for IgG (FcyR), such as FcyRI (CD64), FcyRII (CD32) and FcyRIII (CD16) so as to focus cellular defense mechanisms to the cell expressing the particular antigen. Bispecific antibodies can also be used to direct cytotoxic agents to cells which express a particular antigen. These antibodies possess an antigen-binding arm and an arm which binds a cytotoxic agent or a radionuclide chelator, such as EOTUBE, DPTA, DOTA, or TETA. Another bispecific antibody of interest binds the protein antigen described herein and further binds tissue factor (TF).

4.13.7 HETEROCONJUGATE ANTIBODIES

Heteroconjugate antibodies are also within the scope of the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells (U.S. Patent No. 4,676,980), and for treatment of HIV infection (WO 91/00360; WO 92/200373; EP 03089). It is contemplated that the antibodies can be prepared in vitro using known methods in synthetic protein chemistry, including those involving crosslinking agents. For example, immunotoxins can be constructed using a disulfide exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-mercaptobutyrimidate and those disclosed, for example, in U.S. Patent No. 4,676,980.

4.13.8 EFFECTOR FUNCTION ENGINEERING

It can be desirable to modify the antibody of the invention with respect to effector function, so as to enhance, e.g., the effectiveness of the antibody in treating cancer. For example, cysteine

residue(s) can be introduced into the Fc region, thereby allowing interchain disulfide bond formation in this region. The homodimeric antibody thus generated can have improved internalization capability and/or increased complement-mediated cell killing and antibody-dependent cellular cytotoxicity (ADCC). See Caron et al., J. Exp Med., 176: 1191-1195 (1992) and Shopes, J. Immunol., 148: 2918-2922 (1992). Homodimeric antibodies with enhanced antitumor activity can also be prepared using heterobifunctional cross-linkers as described in Wolff et al. Cancer Research, 53: 2560-2565 (1993). Alternatively, an antibody can be engineered that has dual Fc regions and can thereby have enhanced complement lysis and ADCC capabilities. See Stevenson et al., Anti-Cancer Drug Design, 3: 219-230 (1989).

4.13.9 IMMUNOCONJUGATES

The invention also pertains to immunoconjugates comprising an antibody conjugated to a cytotoxic agent such as a chemotherapeutic agent, toxin (e.g., an enzymatically active toxin of bacterial, fungal, plant, or animal origin, or fragments thereof), or a radioactive isotope (i.e., a radioconjugate).

Chemotherapeutic agents useful in the generation of such immunoconjugates have been described above. Enzymatically active toxins and fragments thereof that can be used include diphtheria A chain, nonbinding active fragments of diphtheria toxin, exotoxin A chain (from Pseudomonas aeruginosa), ricin A chain, abrin A chain, modeccin A chain, alpha-sarcin, Aleurites fordii proteins, dianthin proteins, Phytolaca americana proteins (PAPI, PAPII, and PAP-S), momordica charantia inhibitor, curcin, crotin, sapaonaria officinalis inhibitor, gelonin, mitogellin, restrictocin, phenomycin, enomycin, and the tricothecenes. A variety of radionuclides are available for the production of radioconjugated antibodies. Examples include ²¹²Bi, ¹³¹I, ¹³¹In, ⁹⁰Y, and ¹⁸⁶Re.

Conjugates of the antibody and cytotoxic agent are made using a variety of bifunctional protein-coupling agents such as N-succinimidyl-3-(2-pyridyldithiol) propionate (SPDP), iminothiolane (IT), bifunctional derivatives of imidoesters (such as dimethyl adipimidate HCL), active esters (such as disuccinimidyl suberate), aldehydes (such as glutareldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as tolyene 2,6-diisocyanate), and bis-active fluorine compounds (such as 1,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin can be prepared as described in Vitetta et al., Science, 238: 1098 (1987). Carbon-14-labeled 1-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX-DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. See WO94/11026.

In another embodiment, the antibody can be conjugated to a "receptor" (such as streptavidin) for utilization in tumor pretargeting wherein the antibody-receptor conjugate is administered to the patient, followed by removal of unbound conjugate from the circulation using a clearing agent and then administration of a "ligand" (e.g., avidin) that is in turn conjugated to a cytotoxic agent.

4.14 COMPUTER READABLE SEQUENCES

In one application of this embodiment, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer readable media" refers to any medium which can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention. As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can be represented in a word processing text file, formatted in commercially-available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g. text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

By providing any of the nucleotide sequences SEQ ID NO: 1-444 or a representative fragment thereof; or a nucleotide sequence at least 95% identical to any of the nucleotide sequences of SEQ ID NO: 1-444 in computer readable form, a skilled artisan can routinely access the sequence information for a variety of purposes. Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer

readable medium. The examples which follow demonstrate how software which implements the BLAST (Altschul et al., J. Mol. Biol. 215:403-410 (1990)) and BLAZE (Brutlag et al., Comp. Chem. 17:203-207 (1993)) search algorithms on a Sybase system is used to identify open reading frames (ORFs) within a nucleic acid sequence. Such ORFs may be protein encoding fragments and may be useful in producing commercially important proteins such as enzymes used in fermentation reactions and in the production of commercially useful metabolites.

As used herein, "a computer-based system" refers to the hardware means, software means, and data storage means used to analyze the nucleotide sequence information of the present invention. The minimum hardware means of the computer-based systems of the present invention comprises a central processing unit (CPU), input means, output means, and data storage means. A skilled artisan can readily appreciate that any one of the currently available computer-based systems are suitable for use in the present invention. As stated above, the computer-based systems of the present invention comprise a data storage means having stored therein a nucleotide sequence of the present invention and the necessary hardware means and software means for supporting and implementing a search means. As used herein, "data storage means" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search means" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means. Search means are used to identify fragments or regions of a known sequence which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, Smith-Waterman, MacPattern (EMBL), BLASTN and BLASTA (NPOLYPEPTIDEIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches can be adapted for use in the present computer-based systems. As used herein, a "target sequence" can be any nucleic acid or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. The most preferred sequence length of a target sequence is from about 10 to 300 amino acids, more preferably from about 30 to 100 nucleotide residues. However, it is well recognized that searches for commercially important fragments,

such as sequence fragments involved in gene expression and protein processing, may be of shorter length.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzyme active sites and signal sequences. Nucleic acid target motifs include, but are not limited to, promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

4.15 TRIPLE HELIX FORMATION

In addition, the fragments of the present invention, as broadly described, can be used to control gene expression through triple helix formation or antisense DNA or RNA, both of which methods are based on the binding of a polynucleotide sequence to DNA or RNA.

Polynucleotides suitable for use in these methods are preferably 20 to 40 bases in length and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 15241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Olmno, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide.

4.16 DIAGNOSTIC ASSAYS AND KITS

The present invention further provides methods to identify the presence or expression of one of the ORFs of the present invention, or homolog thereof, in a test sample, using a nucleic acid probe or antibodies of the present invention, optionally conjugated or otherwise associated with a suitable label.

In general, methods for detecting a polynucleotide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polynucleotide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polynucleotide of the invention is detected in the sample. Such methods can also comprise contacting a sample under stringent hybridization conditions with nucleic acid primers

that anneal to a polynucleotide of the invention under such conditions, and amplifying annealed polynucleotides, so that if a polynucleotide is amplified, a polynucleotide of the invention is detected in the sample.

In general, methods for detecting a polypeptide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polypeptide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polypeptide of the invention is detected in the sample.

In detail, such methods comprise incubating a test sample with one or more of the antibodies or one or more of the nucleic acid probes of the present invention and assaying for binding of the nucleic acid probes or antibodies to components within the test sample.

Conditions for incubating a nucleic acid probe or antibody with a test sample vary. Incubation conditions depend on the format employed in the assay, the detection methods employed, and the type and nature of the nucleic acid probe or antibody used in the assay. One skilled in the art will recognize that any one of the commonly available hybridization, amplification or immunological assay formats can readily be adapted to employ the nucleic acid probes or antibodies of the present invention. Examples of such assays can be found in Chard, T., An Introduction to Radioimmunoassay and Related Techniques, Elsevier Science Publishers, Amsterdam, The Netherlands (1986); Bullock, G.R. et al., Techniques in Immunocytochemistry, Academic Press, Orlando, FL Vol. 1 (1982), Vol. 2 (1983), Vol. 3 (1985); Tijssen, P., Practice and Theory of immunoassays: Laboratory Techniques in Biochemistry and Molecular Biology, Elsevier Science Publishers, Amsterdam, The Netherlands (1985). The test samples of the present invention include cells, protein or membrane extracts of cells, or biological fluids such as sputum, blood, serum, plasma, or urine. The test sample used in the above-described method will vary based on the assay format, nature of the detection method and the tissues, cells or extracts used as the sample to be assayed. Methods for preparing protein extracts or membrane extracts of cells are well known in the art and can be readily be adapted in order to obtain a sample which is compatible with the system utilized.

In another embodiment of the present invention, kits are provided which contain the necessary reagents to carry out the assays of the present invention. Specifically, the invention provides a compartment kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the probes or antibodies of the present invention; and (b) one or more other containers comprising one or more of the following: wash reagents, reagents capable of detecting presence of a bound probe or antibody.

In detail, a compartment kit includes any kit in which reagents are contained in separate containers. Such containers include small glass containers, plastic containers or strips of plastic

or paper. Such containers allows one to efficiently transfer reagents from one compartment to another compartment such that the samples and reagents are not cross-contaminated, and the agents or solutions of each container can be added in a quantitative fashion from one compartment to another. Such containers will include a container which will accept the test sample, a container which contains the antibodies used in the assay, containers which contain wash reagents (such as phosphate buffered saline, Tris-buffers, etc.), and containers which contain the reagents used to detect the bound antibody or probe. Types of detection reagents include labeled nucleic acid probes, labeled secondary antibodies, or in the alternative, if the primary antibody is labeled, the enzymatic, or antibody binding reagents which are capable of reacting with the labeled antibody. One skilled in the art will readily recognize that the disclosed probes and antibodies of the present invention can be readily incorporated into one of the established kit formats which are well known in the art.

4.17 MEDICAL IMAGING

The novel polypeptides and binding partners of the invention are useful in medical imaging of sites expressing the molecules of the invention (e.g., where the polypeptide of the invention is involved in the immune response, for imaging sites of inflammation or infection). See, e.g., Kunkel et al., U.S. Pat. NO. 5,413,778. Such methods involve chemical attachment of a labeling or imaging agent, administration of the labeled polypeptide to a subject in a pharmaceutically acceptable carrier, and imaging the labeled polypeptide *in vivo* at the target site.

4.18 SCREENING ASSAYS

Using the isolated proteins and polynucleotides of the invention, the present invention further provides methods of obtaining and identifying agents which bind to a polypeptide encoded by an ORF corresponding to any of the nucleotide sequences set forth in SEQ ID NO: 1-444, or bind to a specific domain of the polypeptide encoded by the nucleic acid. In detail, said method comprises the steps of:

- (a) contacting an agent with an isolated protein encoded by an ORF of the present invention, or nucleic acid of the invention; and
 - (b) determining whether the agent binds to said protein or said nucleic acid.

In general, therefore, such methods for identifying compounds that bind to a polynucleotide of the invention can comprise contacting a compound with a polynucleotide of the invention for a time sufficient to form a polynucleotide/compound complex, and detecting

the complex, so that if a polynucleotide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Likewise, in general, therefore, such methods for identifying compounds that bind to a polypeptide of the invention can comprise contacting a compound with a polypeptide of the invention for a time sufficient to form a polypeptide/compound complex, and detecting the complex, so that if a polypeptide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Methods for identifying compounds that bind to a polypeptide of the invention can also comprise contacting a compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a receptor gene sequence in the cell, and detecting the complex by detecting reporter gene sequence expression, so that if a polypeptide/compound complex is detected, a compound that binds a polypeptide of the invention is identified.

Compounds identified via such methods can include compounds which modulate the activity of a polypeptide of the invention (that is, increase or decrease its activity, relative to activity observed in the absence of the compound). Alternatively, compounds identified via such methods can include compounds which modulate the expression of a polynucleotide of the invention (that is, increase or decrease expression relative to expression levels observed in the absence of the compound). Compounds, such as compounds identified via the methods of the invention, can be tested using standard assays well known to those of skill in the art for their ability to modulate activity/expression.

The agents screened in the above assay can be, but are not limited to, peptides, carbohydrates, vitamin derivatives, or other pharmaceutical agents. The agents can be selected and screened at random or rationally selected or designed using protein modeling techniques.

For random screening, agents such as peptides, carbohydrates, pharmaceutical agents and the like are selected at random and are assayed for their ability to bind to the protein encoded by the ORF of the present invention. Alternatively, agents may be rationally selected or designed. As used herein, an agent is said to be "rationally selected or designed" when the agent is chosen based on the configuration of the particular protein. For example, one skilled in the art can readily adapt currently available procedures to generate peptides, pharmaceutical agents and the like, capable of binding to a specific peptide sequence, in order to generate rationally designed antipeptide peptides, for example see Hurby et al., Application of Synthetic Peptides: Antisense Peptides," In Synthetic Peptides, A User's Guide, W.H. Freeman, NY (1992), pp. 289-307, and Kaspczak et al., Biochemistry 28:9230-8 (1989), or pharmaceutical agents, or the like.

In addition to the foregoing, one class of agents of the present invention, as broadly described, can be used to control gene expression through binding to one of the ORFs or EMFs of the present invention. As described above, such agents can be randomly screened or rationally designed/selected. Targeting the ORF or EMF allows a skilled artisan to design sequence specific or element specific agents, modulating the expression of either a single ORF or multiple ORFs which rely on the same EMF for expression control. One class of DNA binding agents are agents which contain base residues which hybridize or form a triple helix formation by binding to DNA or RNA. Such agents can be based on the classic phosphodiester, ribonucleic acid backbone, or can be a variety of sulfhydryl or polymeric derivatives which have base attachment capacity.

Agents suitable for use in these methods preferably contain 20 to 40 bases and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide and other DNA binding agents.

Agents which bind to a protein encoded by one of the ORFs of the present invention can be used as a diagnostic agent. Agents which bind to a protein encoded by one of the ORFs of the present invention can be formulated using known techniques to generate a pharmaceutical composition.

4.19 USE OF NUCLEIC ACIDS AS PROBES

Another aspect of the subject invention is to provide for polypeptide-specific nucleic acid hybridization probes capable of hybridizing with naturally occurring nucleotide sequences. The hybridization probes of the subject invention may be derived from any of the nucleotide sequences SEQ ID NO: 1-444. Because the corresponding gene is only expressed in a limited number of tissues, a hybridization probe derived from any of the nucleotide sequences SEQ ID NO: 1-444 can be used as an indicator of the presence of RNA of cell type of such a tissue in a sample.

Any suitable hybridization technique can be employed, such as, for example, in situ hybridization. PCR as described in US Patents Nos. 4,683,195 and 4,965,188 provides

additional uses for oligonucleotides based upon the nucleotide sequences. Such probes used in PCR may be of recombinant origin, may be chemically synthesized, or a mixture of both. The probe will comprise a discrete nucleotide sequence for the detection of identical sequences or a degenerate pool of possible sequences for identification of closely related genomic sequences.

Other means for producing specific hybridization probes for nucleic acids include the cloning of nucleic acid sequences into vectors for the production of mRNA probes. Such vectors are known in the art and are commercially available and may be used to synthesize RNA probes in vitro by means of the addition of the appropriate RNA polymerase as T7 or SP6 RNA polymerase and the appropriate radioactively labeled nucleotides. The nucleotide sequences may be used to construct hybridization probes for mapping their respective genomic sequences. The nucleotide sequence provided herein may be mapped to a chromosome or specific regions of a chromosome using well known genetic and/or chromosomal mapping techniques. These techniques include in situ hybridization, linkage analysis against known chromosomal markers, hybridization screening with libraries or flow-sorted chromosomal preparations specific to known chromosomes, and the like. The technique of fluorescent in situ hybridization of chromosome spreads has been described, among other places, in Verma et al (1988) Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York NY.

Fluorescent in situ hybridization of chromosomal preparations and other physical chromosome mapping techniques may be correlated with additional genetic map data. Examples of genetic map data can be found in the 1994 Genome Issue of Science (265:1981f). Correlation between the location of a nucleic acid on a physical chromosomal map and a specific disease (or predisposition to a specific disease) may help delimit the region of DNA associated with that genetic disease. The nucleotide sequences of the subject invention may be used to detect differences in gene sequences between normal, carrier or affected individuals.

4.20 PREPARATION OF SUPPORT BOUND OLIGONUCLEOTIDES

Oligonucleotides, i.e., small nucleic acid segments, may be readily prepared by, for example, directly synthesizing the oligonucleotide by chemical means, as is commonly practiced using an automated oligonucleotide synthesizer.

Support bound oligonucleotides may be prepared by any of the methods known to those of skill in the art using any suitable support such as glass, polystyrene or Teflon. One strategy is to precisely spot oligonucleotides synthesized by standard synthesizers. Immobilization can be achieved using passive adsorption (Inouye & Hondo, (1990) J. Clin. Microbiol. 28(6) 1469-72); using UV light (Nagata et al., 1985; Dahlen et al., 1987; Morrissey & Collins, (1989) Mol. Cell

Probes 3(2) 189-207) or by covalent binding of base modified DNA (Keller *et al.*, 1988; 1989); all references being specifically incorporated herein.

Another strategy that may be employed is the use of the strong biotin-streptavidin interaction as a linker. For example, Broude *et al.* (1994) Proc. Natl. Acad. Sci. USA 91(8) 3072-6, describe the use of biotinylated probes, although these are duplex probes, that are immobilized on streptavidin-coated magnetic beads. Streptavidin-coated beads may be purchased from Dynal, Oslo. Of course, this same linking chemistry is applicable to coating any surface with streptavidin. Biotinylated probes may be purchased from various sources, such as, e.g., Operon Technologies (Alameda, CA).

Nunc Laboratories (Naperville, IL) is also selling suitable material that could be used. Nunc Laboratories have developed a method by which DNA can be covalently bound to the microwell surface termed Covalink NH. CovaLink NH is a polystyrene surface grafted with secondary amino groups (>NH) that serve as bridge-heads for further covalent coupling. CovaLink Modules may be purchased from Nunc Laboratories. DNA molecules may be bound to CovaLink exclusively at the 5'-end by a phosphoramidate bond, allowing immobilization of more than 1 pmol of DNA (Rasmussen et al., (1991) Anal. Biochem. 198(1) 138-42).

The use of CovaLink NH strips for covalent binding of DNA molecules at the 5'-end has been described (Rasmussen et al., (1991). In this technology, a phosphoramidate bond is employed (Chu et al., (1983) Nucleic Acids Res. 11(8) 6513-29). This is beneficial as immobilization using only a single covalent bond is preferred. The phosphoramidate bond joins the DNA to the CovaLink NH secondary amino groups that are positioned at the end of spacer arms covalently grafted onto the polystyrene surface through a 2 nm long spacer arm. To link an oligonucleotide to CovaLink NH via an phosphoramidate bond, the oligonucleotide terminus must have a 5'-end phosphate group. It is, perhaps, even possible for biotin to be covalently bound to CovaLink and then streptavidin used to bind the probes.

More specifically, the linkage method includes dissolving DNA in water (7.5 ng/ μ l) and denaturing for 10 min. at 95°C and cooling on ice for 10 min. Ice-cold 0.1 M 1-methylimidazole, pH 7.0 (1-MeIm₇), is then added to a final concentration of 10 mM 1-MeIm₇. The single-stranded DNA solution is then dispensed into CovaLink NH strips (75 μ l/well) standing on ice.

Carbodiimide 0.2 M 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC), dissolved in 10 mM 1-MeIm₇, is made fresh and 25 µl added per well. The strips are incubated for 5 hours at 50°C. After incubation the strips are washed using, e.g., Nunc-Immuno Wash; first the wells are washed 3 times, then they are soaked with washing solution for 5 min., and finally they are washed 3 times (where in the washing solution is 0.4 N NaOH, 0.25% SDS heated to 50°C).

It is contemplated that a further suitable method for use with the present invention is that described in PCT Patent Application WO 90/03382 (Southern & Maskos), incorporated herein by reference. This method of preparing an oligonucleotide bound to a support involves attaching a nucleoside 3'-reagent through the phosphate group by a covalent phosphodiester link to aliphatic hydroxyl groups carried by the support. The oligonucleotide is then synthesized on the supported nucleoside and protecting groups removed from the synthetic oligonucleotide chain under standard conditions that do not cleave the oligonucleotide from the support. Suitable reagents include nucleoside phosphoramidite and nucleoside hydrogen phosphorate.

An on-chip strategy for the preparation of DNA probe for the preparation of DNA probe arrays may be employed. For example, addressable laser-activated photodeprotection may be employed in the chemical synthesis of oligonucleotides directly on a glass surface, as described by Fodor *et al.* (1991) Science 251(4995) 767-73, incorporated herein by reference. Probes may also be immobilized on nylon supports as described by Van Ness *et al.* (1991) Nucleic Acids Res. 19(12) 3345-50; or linked to Teflon using the method of Duncan & Cavalier (1988) Anal. Biochem. 169(1) 104-8; all references being specifically incorporated herein.

To link an oligonucleotide to a nylon support, as described by Van Ness *et al.* (1991), requires activation of the nylon surface via alkylation and selective activation of the 5'-amine of oligonucleotides with cyanuric chloride.

One particular way to prepare support bound oligonucleotides is to utilize the light-generated synthesis described by Pease *et al.*, (1994) PNAS USA 91(11) 5022-6, incorporated herein by reference). These authors used current photolithographic techniques to generate arrays of immobilized oligonucleotide probes (DNA chips). These methods, in which light is used to direct the synthesis of oligonucleotide probes in high-density, miniaturized arrays, utilize photolabile 5'-protected *N*-acyl-deoxynucleoside phosphoramidites, surface linker chemistry and versatile combinatorial synthesis strategies. A matrix of 256 spatially defined oligonucleotide probes may be generated in this manner.

4.21 PREPARATION OF NUCLEIC ACID FRAGMENTS

The nucleic acids may be obtained from any appropriate source, such as cDNAs, genomic DNA, chromosomal DNA, microdissected chromosome bands, cosmid or YAC inserts, and RNA, including mRNA without any amplification steps. For example, Sambrook *et al.* (1989) describes three protocols for the isolation of high molecular weight DNA from mammalian cells (p. 9.14-9.23).

DNA fragments may be prepared as clones in M13, plasmid or lambda vectors and/or prepared directly from genomic DNA or cDNA by PCR or other amplification methods. Samples

may be prepared or dispensed in multiwell plates. About 100-1000 ng of DNA samples may be prepared in 2-500 ml of final volume.

The nucleic acids would then be fragmented by any of the methods known to those of skill in the art including, for example, using restriction enzymes as described at 9.24-9.28 of Sambrook *et al.* (1989), shearing by ultrasound and NaOH treatment.

Low pressure shearing is also appropriate, as described by Schriefer *et al.* (1990) Nucleic Acids Res. 18(24) 7455-6, incorporated herein by reference). In this method, DNA samples are passed through a small French pressure cell at a variety of low to intermediate pressures. A lever device allows controlled application of low to intermediate pressures to the cell. The results of these studies indicate that low-pressure shearing is a useful alternative to sonic and enzymatic DNA fragmentation methods.

One particularly suitable way for fragmenting DNA is contemplated to be that using the two base recognition endonuclease, *CviJI*, described by Fitzgerald *et al.* (1992) Nucleic Acids Res. 20(14) 3753-62. These authors described an approach for the rapid fragmentation and fractionation of DNA into particular sizes that they contemplated to be suitable for shotgun cloning and sequencing.

The restriction endonuclease CviJI normally cleaves the recognition sequence PuGCPy between the G and C to leave blunt ends. Atypical reaction conditions, which alter the specificity of this enzyme (CviJI**), yield a quasi-random distribution of DNA fragments form the small molecule pUC19 (2688 base pairs). Fitzgerald et al. (1992) quantitatively evaluated the randomness of this fragmentation strategy, using a CviJI** digest of pUC19 that was size fractionated by a rapid gel filtration method and directly ligated, without end repair, to a lac Z minus M13 cloning vector. Sequence analysis of 76 clones showed that CviJI** restricts pyGCPy and PuGCPu, in addition to PuGCPy sites, and that new sequence data is accumulated at a rate consistent with random fragmentation.

As reported in the literature, advantages of this approach compared to sonication and agarose gel fractionation include: smaller amounts of DNA are required (0.2-0.5 µg instead of 2-5 µg); and fewer steps are involved (no preligation, end repair, chemical extraction, or agarose gel electrophoresis and elution are needed

Irrespective of the manner in which the nucleic acid fragments are obtained or prepared, it is important to denature the DNA to give single stranded pieces available for hybridization. This is achieved by incubating the DNA solution for 2-5 minutes at 80-90°C. The solution is then cooled quickly to 2°C to prevent renaturation of the DNA fragments before they are contacted with the chip. Phosphate groups must also be removed from genomic DNA by methods known in the art.

4.22 PREPARATION OF DNA ARRAYS

Arrays may be prepared by spotting DNA samples on a support such as a nylon membrane. Spotting may be performed by using arrays of metal pins (the positions of which correspond to an array of wells in a microtiter plate) to repeated by transfer of about 20 nl of a DNA solution to a nylon membrane. By offset printing, a density of dots higher than the density of the wells is achieved. One to 25 dots may be accommodated in 1 mm², depending on the type of label used. By avoiding spotting in some preselected number of rows and columns, separate subsets (subarrays) may be formed. Samples in one subarray may be the same genomic segment of DNA (or the same gene) from different individuals, or may be different, overlapped genomic clones. Each of the subarrays may represent replica spotting of the same samples. In one example, a selected gene segment may be amplified from 64 patients. For each patient, the amplified gene segment may be in one 96-well plate (all 96 wells containing the same sample). A plate for each of the 64 patients is prepared. By using a 96-pin device, all samples may be spotted on one 8 x 12 cm membrane. Subarrays may contain 64 samples, one from each patient. Where the 96 subarrays are identical, the dot span may be 1 mm² and there may be a 1 mm space between subarrays.

Another approach is to use membranes or plates (available from NUNC, Naperville, Illinois) which may be partitioned by physical spacers e.g. a plastic grid molded over the membrane, the grid being similar to the sort of membrane applied to the bottom of multiwell plates, or hydrophobic strips. A fixed physical spacer is not preferred for imaging by exposure to flat phosphor-storage screens or x-ray films.

The present invention is illustrated in the following examples. Upon consideration of the present disclosure, one of skill in the art will appreciate that many other embodiments and variations may be made in the scope of the present invention. Accordingly, it is intended that the broader aspects of the present invention not be limited to the disclosure of the following examples. The present invention is not to be limited in scope by the exemplified embodiments which are intended as illustrations of single aspects of the invention, and compositions and methods which are functionally equivalent are within the scope of the invention. Indeed, numerous modifications and variations in the practice of the invention are expected to occur to those skilled in the art upon consideration of the present preferred embodiments. Consequently, the only limitations which should be placed upon the scope of the invention are those which appear in the appended claims.

All references cited within the body of the instant specification are hereby incorporated by reference in their entirety.

5. EXAMPLES

5.1 EXAMPLE 1

Novel Nucleic Acid Sequences Obtained From Various Libraries

A plurality of novel nucleic acids were obtained from cDNA libraries prepared from various human tissues and in some cases isolated from a genomic library derived from human chromosome using standard PCR, SBH sequence signature analysis and Sanger sequencing techniques. The inserts of the library were amplified with PCR using primers specific for the vector sequences which flank the inserts. Clones from cDNA libraries were spotted on nylon membrane filters and screened with oligonucleotide probes (e.g., 7-mers) to obtain signature sequences. The clones were clustered into groups of similar or identical sequences. Representative clones were selected for sequencing.

In some cases, the 5' sequence of the amplified inserts was then deduced using a typical Sanger sequencing protocol. PCR products were purified and subjected to fluorescent dye terminator cycle sequencing. Single pass gel sequencing was done using a 377 Applied Biosystems (ABI) sequencer to obtain the novel nucleic acid sequences

5.2 EXAMPLE 2

Assemblage of Novel Nucleic Acids

The nucleic acids of the present invention, designated as SEQ ID NO: 1-444 were assembled using an EST sequence as a seed. Then a recursive algorithm was used to extend the seed EST into an extended assemblage, by pulling additional sequences from different databases (i.e., Hyseq's database containing EST sequences, dbEST, gb pri, and UniGene, and exons from public domain genomic sequences predicated by GenScan) that belong to this assemblage. The algorithm terminated when there was no additional sequences from the above databases that would extend the assemblage. Further, inclusion of component sequences into the assemblage was based on a BLASTN hit to the extending assemblage with BLAST score greater than 300 and percent identity greater than 95%.

Using PHRAP (Univ. of Washington) or CAP4 (Paracel), full-length gene sequences and their corresponding protein sequences were generated from the assemblage. Any frame shifts and incorrect stop codons were corrected by hand editing. During editing, the sequence was checked using FASTXY algorithm against Genbank (i.e., dbEST, gb pri, UniGene, and Genpept). Other computer programs which may have been used in the editing process were phredPhrap and Consed (University of Washington) and ed-ready, ed-ext and gc-zip-2 (Hyseq, Inc.). The full-length nucleotide sequences are shown in the Sequence Listing as SEQ ID NO: 1-444. The corresponding polypeptide sequences are SEQ ID NO: 445-888.

Table 1 shows the various tissue sources of SEQ ID NO: 1-444.

The nearest neighbor results for polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888) were obtained by a BLASTP (version 2.0al 19MP-WashU) search against Genpept release 124 using BLAST algorithm. The nearest neighbor result showed the closest homologue with functional annotation for SEQ ID NO: 1-444 from Genpept. The translated amino acid sequences for which the nucleic acid sequence encodes are shown in the Sequence Listing. The homologues with identifiable functions for SEQ ID NO: 1-444 are shown in Table 2 below.

Using eMatrix software package (Stanford University, Stanford, CA) (Wu et al., J. Comp. Biol., Vol. 6 pp. 219-235 (1999) herein incorporated by reference), polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888) were examined to determine whether they had identifiable signature regions. Table 3 shows the signature region found in the indicated polypeptide sequences, the description of the signature, the eMatrix p-value(s) and the position(s) of the signature within the polypeptide sequence.

Using the pFam software program (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1) pp. 320-322 (1998) herein incorporated by reference) polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888) were examined for domains with homology to certain peptide domains. Table 4 shows the name of the domain found, the description, the p-value and the pFam score for the identified domain within the sequence.

The GeneAtlas™ software package (Molecular Simulations Inc. (MSI), San Diego, CA) was used to predict the three-dimensional structure models for the polypeptides encoded by SEQ ID NO: 1-444 (i.e. SEQ ID NO: 445-888). Models were generated by (1) PSI-BLAST which is a multiple alignment sequence profile-based searching developed by Altschul et al, (Nucl. Acids. Res. 25, 3389-3408 (1997)), (2) High Throughput Modeling (HTM) (Molecular Simulations Inc. (MSI) San Diego, CA,) which is an automated sequence and structure searching procedure (http://www.msi.com/), and (3) SeqFoldTM which is a fold recognition method described by Fischer and Eisenberg (J. Mol. Biol. 209, 779-791 (1998)). This analysis was carried out, in part, by comparing the polypeptides of the invention with the known NMR (nuclear magnetic resonance) and x-ray crystal three-dimensional structures as templates. Table 5 shows, "PDB ID", the Protein DataBase (PDB) identifier given to template structure; "Chain ID", identifier of the subcomponent of the PDB template structure; "Compound Information", information of the PDB template structure and/or its subcomponents; "PDB Function Annotation" gives function of the PDB template as annotated by the PDB files (http://www.rcsb.org/PDB/); start and end amino acid position of the protein sequence aligned; PSI-BLAST score, the verify score, the SeqFold score, and the Potential(s) of Mean Force (PMF). The verify score is produced by GencAtlas™

software (MSI), is based on Dr. Eisenberg's Profile-3D threading program developed in Dr. David Eisenberg's laboratory (US patent no. 5,436,850 and Luthy, Bowie, and Eisenberg, Nature, 356:83-85 (1992)) and a publication by R. Sanchez and A. Sali, Proc. Natl. Acad. Sci. USA, 95:13597-12502. The verify score produced by GeneAtlas normalizes the verify score for proteins with different lengths so that a unified cutoff can be used to select good models as follows:

Verify score (normalized) = (raw score - 1/2 high score)/(1/2 high score)

The PFM score, produced by GeneAtlas™ software (MSI), is a composite scoring function that depends in part on the compactness of the model, sequence identity in the alignment used to build the model, pairwise and surface mean force potentials (MFP). As given in table 8, a verify score between 0 to 1.0, with 1 being the best, represents a good model. Similarly, a PMF score between 0 to 1.0, with 1 being the best, represents a good model. A SeqFold™ score of more than 50 is considered significant. A good model may also be determined by one of skill in the art based all the information in Table 5 taken in totality.

The nucleotide sequence within the sequences that codes for signal peptide sequences and their cleavage sites can be determined from using Neural Network SignalP V1.1 program (from Center for Biological Sequence Analysis, The Technical University of Denmark). The process for identifying prokaryotic and eukaryotic signal peptides and their cleavage sites are also disclosed by Henrik Nielson, Jacob Engelbrecht, Soren Brunak, and Gunnar von Heijne in the publication "Identification of prokaryotic and eukaryotic signal peptides and prediction of their cleavage sites" Protein Engineering, Vol. 10, no. 1, pp. 1-6 (1997), incorporated herein by reference. A maximum S score and a mean S score, as described in the Nielson et al, as reference, were obtained for the polypeptide sequences. Table 6 shows the position of the last amino acid of the signal peptide in each of the polypeptides and the maximum score and mean score associated with that signal peptide.

Table 7 correlates each of SEQ ID NO: 1-444 to a specific chromosomal location.

Table 8 is a correlation table of the novel polynucleotide sequences SEQ ID NO: 1-444, and their corresponding priority nucleotide sequences in the priority application USSN 09/659,671, herein incorporated by reference in its entirety.

TABLE 1

TABLE 1			
Tissue Origin	RNA Source	Library Name	SEQ ID NO:
adult brain	GIBCO	AB3001	4 6-8 12 23 33-34 47 50 55 57-60 62 89 102 104-106 123 144 162 176-177 179 187 194
			248 260 270 279 292 294 297-298 307 322- 323 326 333 336 341 351 450
adult brain	GIBCO	ABD003	6 10 12-15 17-18 26 31 34-35 38-40 42-44
			46 48-50 53 56 59-60 64 66 70-72 80-81
			85-86 98 101 107 116-117 125 130 138-139
			142 144 147 151 160-161 164 173 175-177
			179 184-185 187-188 194-195 198 201 215
			217-218 222 226 228 232 239-240 243-244
			247 252 256 258 260 264-265 267 274-275
			284 288 290 293 298 306-308 314-315 318-
:			320 325-326 333-334 337 341 343 345-346
			351-354 364-365 371 390-391 424-425 429
adult brain	Clontech	ABR001	5 36 43 76 108 128 182-183 212 239 242
<u></u>			260 263 269 296 325 351 364 371-372 423 2 9 11 13 18 23 35 38 42 46 50-51 53-54 60
adult brain	Clontech	ABR006	63 66 85 91 107-108 116-117 120 122 128
			170 178 180 184 187-188 193-194 198 202
1			215 232 243 245 257-258 260 266-267 271
			285 294 301 333-334 337 370 389 394-396
			400 405 412 423 428 434 436 453 458
		ABR008	1 3 7 10-14 16-17 19-23 26-28 34-35 38-39
adult brain	Clontech	ADRUUS	41 43 46-48 51-54 56 60 62 64 66-68 75 82
			86-87 91 96-98 102 104-106 108 110-111
			114 116-118 122 125 127-130 134 138-139
			141-143 145-146 150-151 153 156 158 160
			162 167-170 173-175 177-180 182 185-186
	1		191-194 196-197 200-201 205-206 208-209
			211 213-215 219-220 226-227 231 238 241
			244 246-248 252 256 260 262-265 269 271
	1		273 278-280 282 284 290 292 296 298 301-
	}		302 306 309 311 315-317 322-323 325-327
			329-331 335-337 339 342-343 345-346
			350-355 359-360 362 364 368 370 372 374
		1	376 381 383 385 387 390-395 400-401 405
			410 412 414 417 420-421 423-425 432 440
			447 450-452 459 472-473
adult brain	Clontech	ABR011	174 177 360
adult brain	BioChain	ABR012	334 341
adult brain	BioChain	ABR013	41-42 60 101 163 355
adult brain	Invitrogen	ABR014	53 95 104-106 143 149 177 180 258
adult brain	Invitrogen	ABR015	42 70-72 79 95 112 138-140 163 195 275 288 322-323 341 343 458
adult brain	Invitrogen	ABR016	13 31 60 79 124 136 154 163 333 341 343
			364 370
adult brain	Invitrogen	ABT004	1 11-13 15 18 24-26 34 50 56 68 87 98 104- 106 111 123-124 131-133 137 144 146 173
			189 194 206 224 247-248 260 262 264 269
	ì		272 274 282 298 318 327 335 346 351 356-
			357 372 375 381 392 409-410 421
	Ctrategano	ADP001	2 11-14 24-25 40 42 47 50 52 57-58 69 76
cultured	Stratagene	: ADF001	107 120 144 151 156 163 168 171 194 197
preadipocytes	i	į	199 203-204 215 229 250-251 262 294 333
	I		338 341 415 450 469-473
-4	Cloutsch	ADR002	10-11 16 18 22-23 27-28 33-35 40 43-45 49
adrenal gland	Clontech	ADRUUZ	61 66 85 98 107-108 111 116-117 124 136
			143 145 160 167 173 175 184 187 201 217-

			218 229 249-251 258 262 269 271 273 277
	i		280 287 289 298 301 308 322-323 337 352
			354 360 414 425 445-446 463
adult heart	GIBCO	AHR001	11-13 15 20-23 26-27 30 33-34 37-40 49 53
			56-58 62-65 67-68 76-77 81 86 88 93-94
			101 104-108 112 114 116-117 119 121-125
			128-130 142 144-145 148 150 154-156
			164-165 167 174-176 178-179 182 184
			186-187 189 195 198 200 202 210 213-219
		1	221 228-229 235 238 240 242-243 246-247
			252-253 260 262 264 266-267 269 275 278
			280-281 283 286-289 293-294 297 302-304
			308 311 313 315-316 318-320 322-324
			328-331 333 336 340-341 343 347 355-356
			359 380-382 386-388 413-414 436
adult kidney	GIBCO	AKD001	4-5 8-13 15-18 20-27 33-35 37 39 42 45-46
,			49 52-54 56-59 62 66-67 73 75 77 80-81 83
			85-86 88 91 97-98 100-108 112 117 119
		l	122 124-125 127-129 134 138-142 145-146
			151 153 155 158 160 162 164-165 168 170
			174-176 178-181 186-189 196-199 202 209
			211-212 215-216 222 232 235 237-240 244
	ļ	\	246-248 250-252 257-258 260 262 264
			269-271 275 280 282-284 287-289 291-294
		i	297 303-312 314-317 322-323 325 327-329
			333 336-337 341 357 359 375 403 407-408
		}	413 428 436 469-471
adult kidney	Invitrogen	AKT002	1 10-11 13 17-18 26-27 35 42 54 64 66 73
			77 82 87 91 94 96-97 113-114 118 135 146
			148 160 173-174 182 187 196 198 200 218-
			219 221 239 243-244 249-251 257-258 260
			264 269-270 274-275 284-285 287-288 290
1			302-303 308-309 312 322-324 330 332-333
			335-337 344 346 369 402 404 417 425 428
j			447 462
adult lung	GIBCO	ALG001	4 6 17-18 24-26 39 43-44 46 49 51 53 55
			57-58 76 84 90 95 98 107 111 126 150 155
			157 164 173 176 184 187 195 210 248-249
			252-253 261 264 275 278 281 287 306 309
			312 314 322-323 333 338 340-341 352 358
		}	365 372 403 450
lymph node	Clontech	ALN001	17 19 26 81 85 149 166 218 228 260 264
1			275-276 282 321 333 341 395-396 436
young liver	GIBCO	ALV001	10 12 14 16-18 20 22 33 40 46 48 56 73 82
,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			85 88 100 102 117 126 134 142-143 173
			175 182 197 209 220 237-239 243 246-247
			279 290 294 302 306 309 319-320 327 334
			359
adult liver	Invitrogen	ALV002	10 15 22 24-25 33 49 66 75 86 95 109 111
3441. 11.01		1	130 138-139 148 151 156 187 189 222 237-
			238 246 249 274 282 286 290 298 301 333
	1	İ	371 377 387 395 424-425 430
adult liver	Clontech	ALV003	73 92 294 341
adult ovary	Invitrogen	AOV001	1 4 6-18 23 27-28 30-34 39 42-47 49-50 52-
addit ovary			59 61-62 64-68 75-77 79 81-82 85-88 90
			93-96 98 100-107 109-111 113-115 117
			121 123 125-129 134-136 140 142-145 148
İ			151 154 156 158 160-164 169-171 175-176
			178-179 181-184 187-188 194-196 200
	:		202-204 206 209-210 212 214-215 217-220
Ì			222-224 227 229 233 235 238-240 242-244
l .	ļ		246-253 256 259-260 262 264-265 267 269
!	1	i	1 10 233 230 237 200 202 207 207 207

			·
			275 277-281 283-284 287-294 296-298
	!		301-304 306-308 310-311 313-316 318-326
			328-330 333-342 344 346 349 352-353
			356-358 360-361 363 369 372 375 382-383
			395-396 402-403 405-407 409 415 417-419
			421 425-427 435-436 450 456 467 469-473
adult placenta	Invitrogen	APL001	17 33 157 228 232 264
placenta	Invitrogen	APL002	11-12 23 31 35 51 53 88 104-106 138-139
			151 260 279 296 307 327 334 337
adult spleen	GIBCO	ASP001 .	1 5 11 14 18-20 22-23 33-34 39-40 48-49
			56-59 62 65 87 91 95 98 104-106 108-109
			113-115 124 128 142 145 149-150 158 162-
			163 173 195 198 202 210 224 227 232 250-
			251 253 264 275 277 281 285-286 299 321
			333 399
adult testis	GIBCO	ATS001	8 10-11 13 17 28 43-44 49 54 57-59 62 77
			109 134 138-139 143-144 148 156 163-164
			174 178 182 187-188 200 202 206 216 221
			232 243 258 290 292 294 297-298 304 308-
			309 324 333 336-337 339-341 346 425 472-
			473
adult bladder	Invitrogen	BLD001	11 24-25 44 57-58 65 141 150-151 163 187
		ļ	195 222 224 244-245 247 260 263 277 282
			334 356 369 375 390-391 416 425 472-473
bone marrow	Clontech	BMD001	9 11 13-14 17 19 22-23 27-28 30 32-35 38
			42 46 48-49 51 56-58 62-63 68 74-76 78 80
•			84-85 87 89 91 93 95 97-98 102 104-106
ı			108 111-112 114-115 117 121-122 125
1			138-140 143 146 149 151 154-155 157-171
			174 176 179 184 187 190 195 200 202 205-
			206 210 215 218-222 224-236 239 249 258
	1		262 264 269 285 294 297 304 310 319-321
			324-325 331 333 341-342 363 420 450
bone marrow	Clontech	BMD002	1 5 7 10-11 14 16 19-20 23-25 28 30-31 35
			40 42-43 46 48-49 51 54 56 62 64 68 74 77
			87 89 91 93-96 98-99 104-106 108 110
			114-115 117 121-122 128 138-139 141 143
		1	145-146 149-151 156-158 163 165 169-171
			174 178-179 181 195 199 203-206 210 212
			216 218 220-221 224 234-235 240 245 248-
			252 254-255 258-260 262 264 269 276-278
			285 292 302-303 308-309 312 322-324 326
į			333 341 343 346 350 352 355 362 379 384-
			386 394 410 418 420 425 440-441 449-451
			469-474
bone marrow	Clontech	BMD004	97 149
bone marrow	Clontech	BMD007	53 143 149 221
colon	Invitrogen	CLN001	10 34 49 52-53 57-59 145 163 175 197 238
			246-247 315-316 346 364 442
mixture of 16	various vendors	CTL016	10 56
tissues/mRNAs			
mixture of 16	various vendors	CTL021	149
tissues/mRNAs			
mixture of 16	various vendors	CTL028	145 358
tissues/mRNAs			
adult cervix	BioChain	CVX001	1 4 6 11 23-25 27 30 34-35 39-40 46 50 59
			62-64 76-77 80-81 88 93 100 107 111-112
}			116 118-119 122-123 125 136 138-139 143
		1	146 151 155 160 163 176 178-182 184 188
		-	195 209 215 218 221 228 232-233 235 239-
		1	240 250-251 261-262 264 266 278 283 287-
			288 300 306-308 311 319-320 322-323 325
1			

			329 333 337 341 349 361 369 388 397 403
			407 422 425 429-430 435-436 453 469-471
endothelial cells	Stratagene	EDT001	1 4 6-8 10-14 16-28 31 34-35 37 39 42-44
			46 49 51 53-54 56-59 62-63 66 70-72 76-78
			80-81 83-84 88 96 101 104-107 110 114
			116 118-119 122-124 130 134 136 138-140
			143-146 153-155 158 160 162-163 170
			174-175 184 186-187 189 194-195 197 200
		1	203-204 209-210 212 214-217 219 222-223
	•		229 232-233 235-238 240 242 244 246-248
			250-251 253 256 258 260-262 264 267 269-
			270 272 276-277 279-282 284-285 288-292
			294 302-304 308 312 317 319-320 324-325
			329 333-334 336-337 339 341 361 375 378
			384-385 397 404 411 415-416 450 469-473
fetal brain	Clontech	FBR001	46 104-106 175 193 258 341
fetal brain	Clontech	FBR004	193-194 226-227 229 260 264 334
fetal brain	Clontech	FBR006	2-3 9-12 14 16-17 19-20 23 27-28 39 43 52-
Ictal Oldin	0.0		53 59 66-67 76 86 91-92 94-95 97-98 101-
			102 104-108 110-111 116-117 125 127-130
			134 138-139 142-143 145-148 150-151 167
			170 174-175 178-179 185 187 194 198 200-
			201 212 215 226-228 231 241 245 261 264
i			269 276-277 279 281-282 284 290 292 300
	1		302 309 325 327 331 333-334 336-337 345
			350-353 357 362 371 376 382-383 388-389
		İ	392 395-396 399 401 414 420 427 432 453
			456 458 472-473
fetal brain	Clontech	FBRS03	199 291 402
fetal brain	Invitrogen	FBT002	12 15 19-20 23 30-31 47 53 59 76 86 94 99
Tetal Olain	Invitroge.		104-107 128 131-133 136 144 148 150 163
		1	168 173 175 185 201 220 233 244 250-251
			254-255 262 273 282 307 309 315-316 322-
			323 327 334-335 337 341 351 364 375 392
			409-412 421 423 469-473
fetal heart	Invitrogen	FHR001	34 43 81 87 129 134 138-139 145 200 288-
Tetal fleat	nivia ogon		289 304 315-316 392
fetal kidney	Clontech	FKD001	6 8 10 17 50 54 77 86 92 112 114 179 217
letal Kittley	Cionicen		223 241 269 301 319-320 322-323 333 340-
	1	1	341 397-398
fetal kidney	Ciontech	FKD002	141 264 309 341 432
fetal kidney	Invitrogen	FKD007	107 123
	Clontech	FLG001	16 33 92 100 149 257 337 340 396
fetal lung		FLG003	8 13 15 32 39 48 51 56 91-93 130-133 146
fetal lung	Invitrogen	FLG003	148 197-198 222 244 257 262 280 286 294
		İ	302 315-316 327 337 352 364 392 396 440
6 . 11	Clantanh	EI C004	122 209
fetal lung	Clontech	FLG004 FLS001	1-13 15-46 48-51 53 55 57-59 61-95 97-119
fetal liver-spleen	Columbia	FLSOUT	121-136 138-139 141 143 146-155 158 160
	University		162-165 167 169-171 178 184 186-187 189
			195 209 211-212 214 219-221 223 227-229
			234-252 254-265 267-269 272-280 283 288
			290-291 295-299 302 304 307 312-317
		}	322-323 333 341 343-344 352-353 361
			365-367 371 390-391 402 418 441 444 466
1		1	1
		EV 0000	469-473 1-3 5 11-13 15-18 20-22 24-25 27 29-31
fetal liver-spleen	Columbia	FLS002	1-3 3 11-13 13-18 20-22 24-23 21 29-33
	University		33-34 36-40 43-44 46 49 51 53-54 56-59 61
			63-65 67-68 73 75 77 80-81 84-86 88-90
		•	92-95 97-98 100 103 107-109 111 114 119
			121-122 124 126-127 129-134 138-139
			141-146 149 151-153 157-160 162 164-165

			167 169-171 178-180 183 190 193 200 209-
			214 218-219 221-222 225 227 233-235
			238-239 244 246-252 257-261 267 269
			274-275 277 285 288 291 294 296 299 308
			312 314-316 333-334 336 338 341-342
		Ì	352-353 364-366 372 382 395-396 402-403
			405 416 420 425 436 441 443 446 449 457
			469-473
fetal liver-spleen	Columbia	FLS003	1 3 19 40 49 54 57-59 68 73 83 92 100 104-
rear mer spreen	University		106 170 176 183 256 260 279 304 314 386
			398 427 467
fetal liver	Invitrogen	FLV001	2 10 14 18 20 26 35 66 92 124 143 146 151
letal livel	Hividogen	12,000	174-175 186 197 200 209 224 238 244 246
			258 278-279 282 309 333 366 377 400 408
C. III	Clontech	FLV002	92 126 244
fetal liver			34-35 48 51 53 92 104-106 110 128 141
fetal liver	Clontech	FLV004	146 149 151 160 249 292 333 338 341 359
		77. (000)	421 11-12 44 76 110-111 145 148-149 178 187
fetal muscle	Invitrogen	FMS001	226 258 260 262 264-265 290 341-342 400-
	1		
			401 404 456
fetal muscle	Invitrogen	FMS002	8 11 23 46 52 61 102 111 120 130 134 148
			150 170 226 233 292 318 334 355 365 386
			414 418 475
fetal skin	Invitrogen	FSK001	5-6 8 10 12 23 26-29 32-33 40 42 48-50 53
		1	55-56 59 64 67 69-72 76 83 88 98-99 102
			110 113 117 123 127-128 131-133 138-139
		ļ	143 148 160 163 167 174-175 195 209-210
			220-221 223-224 227 237-238 240 244-247
	1		254-255 257-258 262-263 269 277 280-282
			288 290-291 294-295 298-299 307 309
			322-323 327 330 333 336-337 340-341 346
ļ			361 364 375 387 404 408 428 432 434-435
			443 448-449 452 454
fetal skin	Invitrogen	FSK002	9 14 22 34-35 39-40 56 73 104-106 109
lean skiii	Annua og en		142-143 150 160 211 220-221 235 240 249
			259 270 302 330 431 448 450 460
fatal calcan	BioChain	FSP001	276
fetal spleen umbilical cord	BioChain	FUC001	4-6 8 11 13 16 18 21-25 27 32 35 37 39-40
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	Ì		237-239 244-246 249-251 253 257 259 269
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lymphocytes	ATCC	LPC001	7 10-11 14-15 18 20 24-25 27 33 35 43-44
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			114 117 119 123-125 128 130-135 141 143-
			146 148 151 153 156-158 160-164 166-168
			171 174-176 178-179 181 183-184 187-188
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278 292-293 306 329 333 3	
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187 194 224 226 277 298 3	
444	
salivary gland Clontech SAL001 4 7 10 40 66 88 102 104-10	6 126 128 151
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320 348 369	
salivary gland Clontech SALS03 42	
skin fibroblast ATCC SFB001 54	
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small intestine Clontech SIN001 1 9 11 13 15 17-18 22 27 3.	
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thymus	Clontech	THMc02	9-10 15 17 24-25 27-28 34 38 40 43 49 57-58 68 74 77 81 87 94-95 98 104-108 110 115-116 128 136-137 143 146 148-151 158 160 165 197 200 210-211 215 221-222 232 235 241 243 245 248 252 269 278 281 286 288-289 292 302 312 321 325 327 329 331 333 338 345 350 365 378 383 387 412 428 439-440 446 451-452 460 465 469-473
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trachea	Clontech	TRC001	17 23 34 90 93 108 142 151 238 240 246 259 266 333 412 472-473
uterus	Clontech	UTR001	18 20 30-31 50 52 114 125 158 164 168 182 198 206 210 248 254-255 260 273 283 304 311 325 365 383 421 423

The 16 tissue/mRNAs and their vendor sources are as follows: 1) Normal adult brain mRNA (Invitrogen), 2) Normal adult kidney mRNA (Invitrogen), 3) Normal fetal brain mRNA (Invitrogen), 4) Normal adult liver mRNA (Invitrogen), 5) Normal fetal kidney mRNA (Invitrogen), 6) Normal fetal liver mRNA (Invitrogen), 7) normal fetal skin mRNA (Invitrogen), 8) human adrenal gland mRNA (Clontech), 9) Human bone marrow mRNA (Clontech), 10) Human leukemia lymphoblastic mRNA (Clontech), 11) Human thymus mRNA (Clontech), 12) human lymph node mRNA (Clontech), 13) human so\spinal cord mRNA (Clontech), 14) human thyroid mRNA (Clontech), 15) human esophagus mRNA (BioChain), 16) human conceptional umbilical cord mRNA (BioChain).

TABLE 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
445	gi4151328	Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1 alpha mRNA, complete cds.	2344	48
445	gi4151330	Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1 beta mRNA, complete cds.	1694	59
445	gi2555183	Rattus norvegicus	SPA-1 like protein p1294	2324	48
446	gi13517972	Homo sapiens	PR-domain containing protein 17 mRNA, complete cds.	2496	100
446	gi10434545	Homo sapiens	cDNA FLJ12827 fis, clone NT2RP2002939, weakly similar to ZINC FINGER PROTEIN 136.	2496	100
446	gi13623607	Homo sapiens	, zinc finger protein 136 (clone pHZ-20), clone MGC:12711, mRNA, complete cds.	710	42
447	gi6093239	Homo sapiens	mRNA; cDNA DKFZp434O0515 (from clone DKFZp434O0515).	1054	100
447	gi3522970	Homo sapiens	Trio mRNA, complete cds.	216	23
447	AAW27227	Homo sapiens	Human TRIO phosphoprotein.	216	23
448	gi7022890	Homo sapiens	cDNA FLJ10700 fis, clone NT2RP3000665.	2838	96
448	gi10438668	Homo sapiens	cDNA: FLJ22327 fis, clone HRC05572.	1333	100
448	gi7020045	Homo sapiens	cDNA FLJ20140 fis, clone COL07182.	1074	79
449	gi6102903	Homo sapiens	mRNA; cDNA DKFZp566D244 (from clone DKFZp566D244); partial cds.	2601	99
449	gi10434000	Homo sapiens	cDNA FLJ12485 fis, clone NT2RM2000420.	1907	100
449	gi10437387	Homo sapiens	cDNA: FLJ21308 fis, clone COL02131.	1519	69
450	gi7670836	Homo sapiens	hepatocellular carcinoma-associated antigen 66 (HCA66) mRNA, complete cds.	3101	99
450	gi7959764	Homo sapiens	PRO1289	935	100
450	gi927708	Saccharomyce s cerevisiae	Ydr449cp; CAI: 0.18	288	32
451	gi7020902	Homo sapiens	cDNA FLJ20657 fis, clone KAT01069.	3231	99
451	gi11037252	Rattus norvegicus	NPL4	3156	96
451	gi10434779	Homo sapiens	cDNA FLJ12984 fis, clone NT2RP3000047, weakly similar to NPL4 PROTEIN.	2812	99
452	gi13160469	Homo sapiens	WDR13 protein (WDR13) gene, complete cds.	1063	94
452	gi12044400	Homo sapiens	WDR13 protein (WDR13) mRNA, complete cds.	1063	94
452	gi13751862	Mus musculus	WD-repeat protein	1058	93
453	gi12619286	Homo sapiens	mRNA for spinal cord-derived protein FIS8G, complete cds.	1133	100
453	gi7638241	Homo sapiens	mesenchymal stem cell protein DSC92 mRNA, complete cds.	1133	100
453	gi12804543	Homo sapiens	, mesenchymal stem cell protein DSC92, clone MGC:2824, mRNA, complete cds.	1133	100
454	gi13279287	Homo sapiens	, clone IMAGE:3633354, mRNA, partial cds.	2066	100
454	gi5052586	Drosophila melanogaster	BcDNA.GH08385	334	25
454	gi10433073	Homo sapiens	cDNA FLJ11749 fis, clone	190	26

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			HEMBA1005558, weakly similar to NUCLEAR PROTEIN SNF7.		
455	:7010040	Homo sapiens	cDNA FLJ20018 fis, clone ADSE00909.	1698	99
455 455	gi7019840 gi13938166	Homo sapiens	, clone MGC:12617, mRNA, complete cds.	1630	98
455	gi9280376	Homo sapiens	ancient conserved domain protein 3 (ACDP3) mRNA, complete cds.	1271	90
456	gi7020190	Homo sapiens	cDNA FLJ20232 fis, clone COLF5593.	1487	100
456	gi14249896	Homo sapiens	, clone MGC:15774, mRNA, complete cds.	1479	99
456	gi9188416	Homo sapiens	Novel human gene mapping to chomosome 22.	1479	99
457	AAW75093	Homo sapiens	Human secreted protein encoded by gene 37 clone HFVGS85.	369	100
457	gi8895089	Homo sapiens	protein x 013 mRNA, complete cds.	145	41
457	gi14250569	Homo sapiens	, protein x 013, clone MGC:3073, mRNA, complete cds.	145	41
458	gi7020228	Homo sapiens	cDNA FLJ20257 fis, clone COLF7231.	1169	100
458	gi7528184	Drosophila melanogaster	bicoid-interacting protein BIN3	389	45
459	gi11345384	Homo sapiens	vacuolar protein sorting protein 18 (VPS18) mRNA, complete cds.	5102	100
459	AAW48303	Homo sapiens	Amino acid sequence of human deep orange protein.	2555	100
459	gi2832850	Drosophila melanogaster	EG:171E4.1	1316	35
460	gi6966967	Homo sapiens	mRNA for dipeptidyl-peptidase III (DPP3 gene).	3814	99
460	gi13938201	Homo sapiens	, dipeptidylpeptidase III, clone MGC:15061, mRNA, complete cds.	3811	99
460	AAB67571	Homo sapiens	Amino acid sequence of a human hydrolytic enzyme HYENZ3.	3807	99
461	AAY53020	Homo sapiens	Human secreted protein clone qb56_19 protein sequence SEQ ID NO:46.	657	100
461	AAY59788	Homo sapiens	Human normal ovarian tissue derived protein 65.	618	100
461	AAG04028	Homo sapiens	Human secreted protein, SEQ ID NO: 8109.	442	72
462	gi13021843	Homo sapiens	polyadenylate binding protein-interacting protein 2 mRNA, complete cds.	679	100
462	gi12052806	Homo sapiens	mRNA; cDNA DKFZp564F163 (from clone DKFZp564F163); complete cds.	675	99
462	gi7106826	Homo sapiens	HSPC218	673	99
463	gi7023258	Homo sapiens	cDNA FLJ10914 fis, clone OVARC1000212.	1067	100
464	gi7023258	Homo sapiens	cDNA FLJ10914 fis, clone OVARC1000212.	649	72
465	gi7022147	Homo sapiens	cDNA FLJ10233 fis, clone HEMBB1000266.	3464	100
465	gi12224837	Homo sapiens	mRNA; cDNA DKFZp547K202 (from clone DKFZp547K202).	3464	100
465	AAY99662	Homo sapiens	Human GTPase associated protein-13.	3464	100
466	gi7582304	Homo sapiens	BM-016	584	100
466	AAW85610	Homo sapiens	Secreted protein clone eh80_1.	330	97
466	AAW78199	Homo sapiens	Human secreted protein encoded by gene 74 clone HGBAC11.	330	97
467	gi7018410	Homo sapiens	mRNA; cDNA DKFZp566K023 (from clone DKFZp566K023).	1010	100
467	gi9049987	Rattus	X2CR! protein	268	81

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		norvegicus			
468	gi8317213	Homo sapiens	histone acetyltransferase (MOF) mRNA, partial cds.	1625	100
468	gi10433157	Homo sapiens	cDNA FLJ11810 fis, clone HEMBA1006347, moderately similar to MALES-ABSENT ON THE FIRST PROTEIN (EC 2.3.1).	1625	100
468	gi10436400	Homo sapiens	cDNA FLJ14040 fis, clone HEMBA1005513, weakly similar to MALES-ABSENT ON THE FIRST PROTEIN (EC 2.3.1).	1613	99
469	AAY76072	Homo sapiens	Human skin cell protein, SEQ ID NO:327.	668	100
469	AAB56011	Homo sapiens	Skin cell protein, SEQ ID NO: 327.	668	100
470	gi29481	Homo sapiens	Human erythrocyte 2,3- bisphosphoglycerate mutase mRNA EC 2.7.5.4.	1362	100
470	gi 179527	Homo sapiens	Human 2,3-bisphosophoglycerate mutase (BPGM) gene, exon 3.	1362	100
470	AAB11959	Homo sapiens	Glycated human erythrocyte bisphosphoglycerate mutase (BPGM).	.1362	100
471	gi6841472	Homo sapiens	HSPC125	892	100
471	gi12001966	Homo sapiens	clone 015g09 My013 protein mRNA, complete cds.	892	100
471	gi9624483	Homo sapiens	HRPAP20 short form mRNA, complete cds.	640	72
472	gi9367763	Homo sapiens	mRNA for zinc finger protein Cezanne (CEZANNE gene).	2580	100
472	gi6102920	Homo sapiens	mRNA; cDNA DKFZp434H0717 (from clone DKFZp434H0717); partial cds.	2197	100
472	gi7332054	Caenorhabditis elegans	contains similarity to tumor necrosis factors	126	25
473	gi8489813	Homo sapiens	DJ963K23.2 mRNA, complete cds.	1255	100
473	AAB43861	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1306.	1255	100
473	gi9858803	Mus musculus	Zfp228	1090	91
474	gi7020223	Homo sapiens	cDNA FLJ20254 fis, clone COLF6926.	2278	100
474	AAY25743	Homo sapiens	Human secreted protein encoded from gene 33.	917	100
474	AAY76166	Homo sapiens	Human secreted protein encoded by gene 43.	724	94
475	gi14042066	Homo sapiens	cDNA FLJ14503 fis, clone NT2RM1000252, weakly similar to H.sapiens E-MAP-115 mRNA.	159	26
475	g17270600	Arabidopsis thaliana	trichohyalin like protein	156	25
475	g1180195	Homo sapiens	Human aorta caldesmon mRNA, complete cds.	145	25
476	g111066250	Homo sapiens	presenilins associated rhomboid-like protein (PARL) mRNA, complete cds.	2030	100
476	gi13177766	Homo sapiens	, Similar to presentiins associated rhomboid-like protein, clone MGC:4756, mRNA, complete cds.	1107	99
476	g17959883	Homo sapiens	PRO2207	986	100
477	AAY91941	Homo sapiens	Human chaperone protein 2 (HCHP-2).	1977	100
477	gi7019854	Homo sapiens	cDNA FLJ20027 fis, clone ADSE01901.	1965	99
477	gi6567172	Mus musculus	mDj10	1863	93
478	gi13937971	Homo sapiens	; Similar to RIKEN cDNA 1110005A23 gene, clone MGC:14726, mRNA,	1040	100

SEQ ID	Accession	Species	Description	Score	% Identity
NO:	No.		complete cds.		lucinity
478	gi13940310	Homo sapiens	HCC-1 gene.	1040	100
478	AAB36609	Homo sapiens	Human FLEXHT-31 protein sequence	1040	100
470	AABSOOOS	i Tomo supiens	SEQ ID NO:31.		
479	gi11065999	Homo sapiens	neuronal calcium binding protein NECAB3 mRNA, complete cds.	1889	99
479	gi10798741	Homo sapiens	XB51 mRNA for X11L-binding protein 51, complete cds.	654	99
479	gi10798743	Mus musculus	X11L binding protein 51	1079	86
480	gi6094684	Homo sapiens	PAC clone RP1-278D1 from X, complete sequence.	3056	92
480	gi10435614	Homo sapiens	cDNA FLJ13568 fis, clone PLACE1008368, weakly similar to RING CANAL PROTEIN.	1847	100
480	gi7023516	Homo sapiens	cDNA FLJ11078 fis, clone PLACE1005102, weakly similar to RING CANAL PROTEIN.	1208	42
481	gi7020424	Homo sapiens	cDNA FLJ20369 fis, clone HEP19364.	2727	100
481	gi1110599	Mus sp.	semaphorin homolog=M-Sema F	2653	86
481	AAB88485	Homo sapiens	Human membrane or secretory protein clone PSEC0078.	1774	100
482	gi4679028	Homo sapiens	HSPC021	1930	100
482	gi5106781	Homo sapiens	HSPC025	1930	100
482	gi12654535	Homo sapiens	, HSPC025, clone MGC:735, mRNA, complete cds.	1930	100
483	gi1145789	Rattus norvegicus	neuroligin 2	4417	98
483	gi7960135	Homo sapiens	neuroligin 3 isoform gene, complete cds, alternatively spliced.	2736	65
483	gi7960131	Homo sapiens	neuroligin 3 isoform HNL3 mRNA, complete cds, alternatively spliced.	2729	65
484	gi14250554	Homo sapiens	, hexokinase 1, clone MGC:1724, mRNA, complete cds.	4725	99
484	gi2873349	Homo sapiens	hexokinase I (HK1) gene, exon 18, complete cds, alternatively spliced.	4725	99
484	gi184021	Homo sapiens	Human hexokinase 1 (HK1) mRNA, complete cds.	4718	99
485	gi8453103	Homo sapiens	zinc finger protein mRNA, complete cds.	3726	100
485	gi13752754	Homo sapiens	zinc finger 1111 mRNA, complete cds.	1689	56
485	gi10436789	Homo sapiens	cDNA FLJ14345 fis, clone THYRO1001189, weakly similar to ZINC FINGER PROTEIN 91.	1683	56
486	AAB56937	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1515.	2341	100
486	gi12804453	Homo sapiens	, Similar to Tu translation elongation factor, mitochondrial, clone MGC:1592, mRNA, complete cds.	2326	100
486	gi899285	Homo sapiens	H.sapiens mRNA for elongations factor Tu-mitochondrial.	2326	100
487	gi9910111	Homo sapiens	myosin X (MYO10) mRNA, complete cds.	10727	99
487	gi6996558	Mus musculus	myosin X	10089	93
487	gi7108753	Homo sapiens	myosin X (MYO10) mRNA, partial cds.	8029	99
488	gi7688687	Homo sapiens	AD-017 protein mRNA, complete cds.	1935	100
488	gi14042251	Homo sapiens	cDNA FLJ14611 fis, clone NT2RP1000988.	1935	100
488	AAY66671	Homo sapiens	Membrane-bound protein PRO1134.	1935	100
489	gi202215	Mus musculus	alpha-tubulin isotype M-alpha-6	2387	100
489	gi14328047	Homo sapiens	, tubulin alpha 4, clone MGC:2379,	2387	100

111

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
110.	1.0.		mRNA, complete cds.		<u> </u>
489	gi1333692	Macaca fascicularis	alpha-tubulin (ATG-initiation codon missing)	2382	100
490	gi5912034	Homo sapiens	mRNA; cDNA DKFZp434N0535 (from clone DKFZp434N0535); partial cds.	6810	99
490	gi5912239	Homo sapiens	mRNA; cDNA DKFZp434O225 (from clone DKFZp434O225); partial cds.	3442	99
490	gi3292939	Drosophila melanogaster	Additional sex combs	295	39
491	gi5912034	Homo sapiens	mRNA; cDNA DKFZp434N0535 (from clone DKFZp434N0535); partial cds.	5941	99
491	gi5912239	Homo sapiens	mRNA; cDNA DKFZp434O225 (from clone DKFZp434O225); partial cds.	2573	99
491	gi3292939	Drosophila melanogaster	Additional sex combs	295	39
492	AAY68778	Homo sapiens	Amino acid sequence of a human phosphorylation effector PHSP-10.	2463	99
492	gi479173	Homo sapiens	H.sapiens nek3 mRNA for protein kinase.	2417	99
492	gi13529320	Mus musculus	Similar to NIMA (never in mitosis gene a)-related expressed kinase 3	1887	73
493	gi13539686	Homo sapiens	protein kinase C and casein kinase substrate 1 (PACSIN1) mRNA, complete cds.	2365	100
493	gi728604	Mus musculus	PACSIN	2250	95
493	gi4324452	Rattus norvegicus	syndapin l	2250	95
494	gi7023749	Homo sapiens	cDNA FLJ11220 fis, clone PLACE1008129.	3994	100
494	gi10433501	Homo sapiens	cDNA FLJ12104 fis, clone HEMBB1002697.	2829	100
494	gi5788108	Homo sapiens	PAC clone RP5-1087M19 from 7q11.23- q21.1, complete sequence.	757	63
495	AAB54375	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:827.	2897	99
495	AAY57923	Homo sapiens	Human transmembrane protein HTMPN-47.	2724	98
495	AAW88628	Homo sapiens	Secreted protein encoded by gene 95 clone HPWAN23.	2686	98
496	gi7959788	Homo sapiens	PRO1635 ·	317	100
496	AAW74852	Homo sapiens	Human secreted protein encoded by gene 124 clone HPCAD23.	143	100
497	gi7707424	Homo sapiens	mRNA for syntaxin 18, complete cds.	1705	100
498	gi1613858	Homo sapiens	Human zinc finger protein zfp47 (zf47) mRNA, partial cds.	1488	83
498	gi13938633	Mus musculus	RIKEN cDNA 2810435N07 gene	1318	60
498	gi9837564	Mus musculus	SCAN-KRAB-zinc finger protein	1242	58
499	AAY27795	Homo sapiens	Human secreted protein encoded by gene No. 79.	1539	99
499	gi10436317	Homo sapiens	cDNA FLJ13986 fis, clone Y79AA1001923, weakly similar to Homo sapiens F-box protein Fbx22 (FBX22) gene.	1370	100
499	gi6164747	Homo sapiens		391	93
500	gi3150052	Homo sapiens	TGF beta receptor associated protein-1 mRNA, complete cds.	4455	100
500	gi14280050	Homo sapiens		382	24
500	gi12718237	Neurospora	related to TGF beta receptor associated	174	31

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		crassa	protein 1		
501	gi7023051	Homo sapiens	cDNA FLJ10796 fis, clone NT2RP4000648, weakly similar to TRANS-ACTING TRANSCRIPTIONAL PROTEIN ICPO.	3360	99
501	gi9651170	Homo sapiens	cell cycle checkpoint protein CHFR mRNA, complete cds.	2491	96
501	AAB20219	Homo sapiens	Human Chfr (checkpoint with FHA and ring finger) protein.	2491	96
502	gi7329074	Homo sapiens	collagen type V alpha 3 chain (COL5A3) mRNA, complete cds.	9671	100
502	gi8568094	Rattus norvegicus	alpha 4 type V collagen	8038	82
502	gi7329072	Mus musculus	collagen type V alpha 3 chain	7970	82
503	gi12654687	Homo sapiens	, clone MGC:2616, mRNA, complete cds.	1161	100
503	gi7769617	Mus musculus	TCE2 ·	1050	89
504	gi12654687	Homo sapiens	, clone MGC:2616, mRNA, complete cds.	1140	96
504	gi7769617	Mus musculus	TCE2	1029	86
505	gi12654687	Homo sapiens	, clone MGC:2616, mRNA, complete cds.	654	100
505	gi7769617	Mus musculus	TCE2	629	92
506	gi14249942	Homo sapiens	, Similar to RIKEN cDNA 0610008P16 gene, clone MGC:15937, mRNA, complete cds.	1609	100
506	AAB56487	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1065.	1167	98
506	gi2828262	Bos taurus	aralkyl acyl-CoA:amino acid N- acyltransferase	597	40
507	gi7688987	Homo sapiens	uncharacterized bone marrow protein BM046	1295	100
507	AAB64387	Home sapiens	Amino acid sequence of human intracellular signalling molecule INTRA19.	1202	94
507	gi9437511	Homo sapiens	BM024	1045	98
508	AAB18979	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1203	100
508	gi6808196	Homo sapiens	mRNA; cDNA DKFZp434P1018 (from clone DKFZp434P1018); partial cds.	938	100
508	gi13960126	Homo sapiens	, Similar to leucine-rich neuronal protein, clone MGC:4126, mRNA, complete cds.	845	100
509	gi13938527	Homo sapiens	, Similar to RIKEN cDNA 2810002N01 gene, clone MGC:2562, mRNA, complete cds.	1048	100
509	AAY35994	Homo sapiens	Extended human secreted protein sequence, SEQ ID NO. 379.	1032	98
509	AAG00345	Homo sapiens	Human secreted protein, SEQ ID NO: 4426.	619	98
510	gi773387	Neurospora crassa	Restriction enzyme inactivation of met-10 complementation in this region. Sequence similarity to S. cerevisiae chromosome VIII cosmid 9205, accession no. U10556 CDS residues 22627-24126	536	35
510	gi487945	Saccharomyce s cerevisiae	Yhr070wp	528	49
510	AAG02508	Homo sapiens	Human secreted protein, SEQ ID NO: 6589.	324	100
511	gi11493195	Homo sapiens	mRNA for LB1 protein.	2614	99
511	gi10434688	Homo sapiens	cDNA FLJ12920 fis, clone NT2RP2004594.	2604	99
511	gi12053201	Homo sapiens	mRNA; cDNA DKFZp434A1031 (from	2604	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			clone DKFZp434A1031); complete cds.		
512	AAW75106	Homo sapiens	Human secreted protein encoded by gene 50 clone HHSDZ57.	471	100
512	AAY59689	Homo sapiens	Secreted protein 26-44-1-B5-CL3_1.	471	100
512	AAY48331	Homo sapiens	Human prostate cancer-associated protein 28.	471	100
514	AAW67888	Homo sapiens	Human secreted protein encoded by gene 82 clone HSKHL65.	921	92
514	gi13436110	Homo sapiens	, Similar to RIKEN cDNA 2310034L04 gene, clone MGC:11061, mRNA, complete cds.	150	28
514	AAY53052	Homo sapiens	Human secreted protein clone df202_3 protein sequence SEQ ID NO:110.	132	33
515	gi7020259	Homo sapiens	cDNA FLJ20276 fis, clone HEP02437.	5378	100
515	gi10432807	Homo sapiens	cDNA FLJ11534 fis, clone HEMBA1002679.	3024	99
515	gi9916	Plasmodium falciparum	liver stage antigen	399	23
516	AAB67448	Homo sapiens	Amino acid sequence of a human chaperone polypeptide.	1190	99
516	gi13477189	Homo sapiens	, Similar to RIKEN cDNA 1300007M11 gene, clone MGC:12943, mRNA, complete cds.	1182	99
516	AAG03527	Homo sapiens	Human secreted protein, SEQ ID NO: 7608.	389	98
517	gi7023782	Homo sapiens	cDNA FLJ11240 fis, clone PLACE1008568.	2796	100
517	AAB08869	Homo sapiens	Amino acid sequence of a human secretory protein.	2792	99
517.	AAB23626	Homo sapiens	Human secreted protein SEQ ID NO: 52.	2792	99
518	gi6460009	Deinococcus radiodurans	citrate lyase, beta subunit	211	30
518	gi 14025765	Mesorhizobiu m loti	citrate lyase beta-subunit	324	31
518	gi14024477	Mesorhizobiu m loti	Citrate lyase beta chain (acyl lyase subunit); CitE	316	33
519	gi14041831	Homo sapiens	cDNA FLJ14357 fis, clone HEMBA1000005, highly similar to DNAJ PROTEIN HOMOLOG MTJ1.	2873	100
519	AAB67447	Homo sapiens	Amino acid sequence of a human chaperone polypeptide.	2481	99
519	gi473847	Mus musculus	dnaJ-like protein	2413	84
520	gi7669968	Homo sapiens	mRNA; cDNA DKFZp761G0313 (from clone DKFZp761G0313).	789	100
520	gi4586315	Homo sapiens	ORCTL3 mRNA for organic-cation transporter like 3, complete cds.	348	38
520	gi4835384	Homo sapiens	DNA, DLEC1 to ORCTL4 gene region, section 1/2 (DLEC1, ORCTL3, ORCTL4 genes, complete cds).	348	38
521	gi7959805	Homo sapiens	PRO0823	344	100
522	gi10434341	Homo sapiens	cDNA FLJ12691 fis, clone NT2RM4002571, weakly similar to H.sapiens mRNA for UDP- GalNAc:polypeptide N- acetylgalactosaminyltransferase (T2).	2605	89
522	gi10436305	Homo sapiens	cDNA FLJ13977 fis, clone Y79AA1001603, weakly similar to POLYPEPTIDE N- ACETYLGALACTOSAMINYLTRANS FERASE (EC 2.4.1.41).	1631	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
522	gi971461	Homo sapiens	H.sapiens mRNA for UDP-	1386	50
			GalNAc:polypeptide N-		
			acetylgalactosaminyltransferase (T2).	}	
523	gi11493500	Homo sapiens	PRO2979	477	100
523	gi38163	Pan	A-gamma-globin	477	100
	8	troglodytes			
523	gi176779	Pan	gamma-2 globin	477	100
	5	troglodytes	3 · · · · · · · · · · · · · · · · · · ·		İ
524	gi5262582	Homo sapiens	mRNA; cDNA DKFZp434K063 (from	3782	99
	3	1	clone DKFZp434K063); partial cds.		
524	gi10438230	Homo sapiens	cDNA: FLJ21993 fis, clone HEP06576.	1416	100
524	AAY21842	Homo sapiens	Human signal peptide-contianing protein	1416	100
			(SIGP) (clone ID 1273453).		
525	gi1928886	Rattus	lin-10 protein homolog	2199	97
	6	norvegicus	, ,	Į	
525	gi10433467	Homo sapiens	cDNA FLJ12076 fis, clone	483	98
			HEMBB 1002442, weakly similar to LIN-		
			10 PROTEIN.		
525	gi5824587	Caenorhabditis	T01G9.2b	668	37
	g	elegans			
526	gi1679607	Mus musculus	myosin-I	4206	84
526	gi1924940	Homo sapiens	H.sapiens mRNA for myosin-IE.	4115	99
526	gi65324	Gallus gallus	brush border myosin IB	3812	76
527	AAB63419	Homo sapiens	Human breast cancer associated antigen	641	99
			protein sequence SEQ ID NO:781.	1	
528	gi13649967	Homo sapiens	fovea-associated SH3 domain binding	558	100
	8		protein (FASH3) mRNA, complete cds.		
528	gi13539561	Homo sapiens	mRNA for SH3BGRL2 protein.	558	100
528	gi5042302	Mus musculus	sh3bgr protein	365	64
529	gi10436540	Homo sapiens	cDNA FLJ14154 fis, clone	1151	99
	8.72		NT2RM1000341.		
529	gi13436011	Mus musculus	RIKEN cDNA 1200013P24 gene	1139	97
529	gi1592161	Methanococcu	ribosomal protein S18 alanine	109	36
	8	s jannaschii	acetyltransferase	'	
530	gi3135314	Homo sapiens	chromosome 7q22 sequence, complete	911	100
	0	•	sequence.		İ
530	gi6752287	Homo sapiens	Novel human gene mapping to	281	51
	٦	· ·	chomosome X.		
531	gi14042818	Homo sapiens	cDNA FLJ14937 fis, clone	2548	97
			PLACE1010231, weakly similar to CELL		
			SURFACE GLYCOPROTEIN EMR1		
			PRECURSOR.		
531	gi2117161	Homo sapiens	H.sapiens mRNA for HE6 Tm7 receptor.	1366	52
531	AAW36903	Homo sapiens	Human epididymis-specific receptor	1366	52
			protein.		
532	gi7417372	Homo sapiens	intracellular hyaluronan-binding protein	2175	99
			mRNA, complete cds.	L	
532	gi7110497	Mus musculus	intracellular hyaluronan-binding protein	1862	85
	1		p57		
532	gi3403154	Homo sapiens	Human Ki-1/57 intracellular antigen	1591	98
	-	•	mRNA, partial cds.	L	<u>L</u>
533	gi10436645	Homo sapiens	cDNA FLJ14235 fis, clone	1585	82
-			NT2RP4000167.	1	
533	gi7020976	Homo sapiens	cDNA FIJ20707 fis, clone KAIA1223.	2195	84
533	gi13276619	Homo sapiens	mRNA; cDNA DKFZp761I0112 (from	1444	99
			clone DKFZp761I0112).	1	
534	gi438880	Rattus	tropomyosin	1186	99
	65000	norvegicus			1
534	gi2978558	Xenopus	alpha-tropomyosin	1089	89

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		laevis			
534	gi438882	Rattus norvegicus	tropomyosin	1086	92
535	gi438880	Rattus norvegicus	tropomyosin	1120	93
535	gi9508585	Homo sapiens	tropomyosin isoform mRNA, complete cds.	1105	93
535	gi12653955	Homo sapiens	, Similar to tropomyosin 4, clone MGC:3261, mRNA, complete cds.	1094	91
536	gi6808111	Homo sapiens	mRNA; cDNA DKFZp434O1230 (from clone DKFZp434O1230); partial cds.	439	100
537	gi6807806	Homo sapiens	mRNA; cDNA DKFZp434K031 (from clone DKFZp434K031); partial cds.	3007	100
537	gi13623334	Homo sapiens	, Similar to DKFZP727C091 protein, clone MGC:10677, mRNA, complete cds.	2392	100
537	AAY25821	Homo sapiens	Human secreted protein fragment encoded from gene 41.	1967	99 .
538	AAB88413	Homo sapiens	Human membrane or secretory protein clone PSEC0170.	1818	99
538	gi6457342	Homo sapiens	E21G4 (E2IG4) mRNA, complete cds.	1813	99
538	AAB24026	Homo sapiens	Human PRO1788 protein sequence SEQ IDNO:18.	1813	99
539	gi6572289	Homo sapiens	mRNA for mitochondrial tryptophanyl-tRNA synthetase (WARS2 gene).	1820	100
539	gi13421159	Caulobacter crescentus	tryptophanyl-tRNA synthetase	727	46
539	gi11992026	Zymomonas mobilis	tryptophany!-tRNA synthase	721	43
540	gi7106630	Homo sapiens	Novel human mRNA from chromosome 1, clone Z98884, has homology to PERIOD CIRCADIAN PROTEIN 3.	6301	99
540	gi13160925	Homo sapiens	mRNA for period (Drosophila) homolog 3 hPER3, complete cds.	6274	99
540	AAB23266	Homo sapiens	Human circadian rhythm protein Per3 (hPer3).	6274	99
541	gi9621744	Homo sapiens	ferritin heavy chain subunit mRNA, complete cds.	968	100
541	gi12654093	Homo sapiens	, ferritin, heavy polypeptide 1, clone MGC:5580, mRNA, complete cds.	968	100
541	gi12655095	Homo sapiens	, ferritin, heavy polypeptide 1, clone MGC:1749, mRNA, complete cds.	968	100
542	gi4902699	Homo sapiens	Novel human gene mapping to chomosome 13.	2372	57
542	gi2341020	Homo sapiens	PAC clone 248O15 from 13q12-q13, complete sequence.	1447	58
542	gil 1907986	Drosophila melanogaster	fry	1054	38
543	gi7582278	Homo sapiens	BM-003	1386	100
543	gi7688983	Homo sapiens	uncharacterized bone marrow protein BM044	1386	100
543	gi1752736	Saccharomyce s cerevisiae	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w	150	35
544	gi1628401	Homo sapiens	H.sapiens mRNA for leucine-rich primary response protein 1.	3936	98
544	gi940821	Rattus norvegicus	LRPR1	2914	73
544	gi2196560	Schizosacchar omyces pombe	Mis6	223	31

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
545	gi7022824	Homo sapiens	cDNA FLJ10656 fis, clone NT2RP2006038.	1574	99
345	gi6841138	Homo sapiens	HSPC099 mRNA, partial cds.	248	36
345	AAG02788	Homo sapiens	Human secreted protein, SEQ ID NO: 6869.	234	85
546	AAB71914	Homo sapiens	Human ISOM-6.	1142	98
546	gi3876969	Caenorhabditis elegans	Similarity to Brugia peptidylprolyl isomerase (TR:G984562), contains similarity to Pfam domain: PF00076 (RNA recognition motif. (a.k.a. RRM, RBD, or RNP domain)), Score=62.0, E-value=4.2e-15, N=1; PF00160 (Cyclophilin type peptidyl-prolyl cis-trans isomerase), Score=78.1, E-value=3.7e-22, N=1	658	52
546	AAG02246	Homo sapiens	Human secreted protein, SEQ ID NO: 6327.	573	100
547	gi603635	Saccharomyce s cerevisiae	Yel044wp	133	25
548	gi5262665	Homo sapiens	mRNA; cDNA DKFZp564B0769 (from clone DKFZp564B0769); partial cds.	1455	99
548	gi6841172	Homo sapiens	HSPC261	716	99
548	gi12803875	Homo sapiens	, Similar to splicing factor, arginine/serine-rich 4, clone MGC:3920, mRNA, complete cds.	352	33
549	gi7582298	Homo sapiens	BM-013	704	100
549	gi9558483	Ciona savignyi	PEM-3	434	55
549	gi1644450	Caenorhabditis elegans	MEX-3	362	65
550	gi4883433	Homo sapiens	mRNA for membrane transport protein (XK gene).	2148	100
550	gi6502963	Mus musculus	KX antigen	1797	81
550	gi2580580	Homo sapiens	testis-specific XK Related Y (XKRY) mRNA, complete cds.	157	31
551	gi7670746	Homo sapiens	UDP-glucose:glycoprotein glucosyltransferase 1 precursor, mRNA, complete cds.	8075	99
551	gi13275621	synthetic construct	Rat RUGT	7371	91
551	gi7677176	Rattus norvegicus	UDP-glucose glycoprotein:glucosyltransferase precursor	7371	91
552	gi7688985	Homo sapiens	uncharacterized bone marrow protein BM045	390	72
553	gi12655091	Homo sapiens	, AD-003 protein, clone MGC:783, mRNA, complete cds.	1177	100
553	gi6523799	Homo sapiens	adrenal gland protein AD-003 mRNA, complete cds.	1168	99
553	gi7105659	Caenorhabditis elegans	contains similarity to Streptomyces peucetius carminomycin 4-O-methyltransferase (GB:L13453)	425	39
554	gi7582282	Homo sapiens	BM-005	3445	99
554	gi7022933	Homo sapiens	cDNA FLJ10725 fis, clone NT2RP3001214.	3312	100
554	gi10435575	Homo sapiens	cDNA FLJ13534 fis, clone PLACE1006445.	1648	100
555	gi12751374	Homo sapiens	paraoxanase-3 mRNA, partial cds.	1819	99
555	gi1333634	Homo sapiens	paraoxonase 3 (PON3) mRNA, 3' end of cds.	1741	98
555	gi12743899	Oryctolagus	рагаохопаѕе 3	1542	82

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		cuniculus			
556	gi7022174	Homo sapiens	cDNA FLJ10252 fis, clone HEMBB1000807.	2826	100
556	gi11596985	Homo sapiens	chromosome 14 clone RP11-361H10 map 14q24.3, complete sequence.	559	36
556	gi7020945	Homo sapiens	cDNA FLJ20689 fis, clone KAIA2890.	510	39
557	gi10434683	Homo sapiens	cDNA FLJ12917 fis, clone	2879	99
	Bresser		NT2RP2004568, weakly similar to PUTATIVE ATP-DEPENDENT RNA HELICASE C30D11.03.		
557	gi13384106	Homo sapiens	RNA helicase-like protein (RHLP) mRNA, complete cds.	2817	99
557	g17020811	Homo sapiens	cDNA FLJ20596 fis, clone KAT08049.	2020	99
558	g:4760710	Brassica rapa	SLL2-S9-protein	284	43
558	g11669601	Arabidopsis thaliana	AR401	280	44
558	gi557805	Saccharomyce s cerevisiae	orf, len: 257, CAI: 0.13	327	34
559	g:13548677	Homo sapiens	MKP-7 mRNA for MAPK phosphatase-7, complete cds.	3418	100
559	g:13990989	Mus musculus	MAP kinase phosphatase-7	3093	90
559	AAB20325	Homo sapiens	Human protein phosphatase and kinase protein-4.	3021	90
560	g110433965	Homo sapiens	cDNA FLJ12464 fis, clone NT2RM1000780.	2196	97
560	gi10434795	Homo sapiens	cDNA FLJ12992 fis, clone NT2RP3000149.	2196	97
560	gi10438048	Homo sapiens	cDNA: FLJ21857 fis, clone HEP02294.	2151	94
561	g110438048	Homo sapiens	cDNA: FLJ21857 fis, clone HEP02294.	2276	97
561	g110433965	Homo sapiens	cDNA FLJ12464 fis, clone NT2RM1000780.	2159	94
561	g110434795	Homo sapiens	cDNA FLJ12992 fis, clone NT2RP3000149.	2159	94
562	g:10433965	Homo sapiens	cDNA FLJ12464 fis, clone NT2RM1000780.	2443	99
562	gi10434795	Homo sapiens	cDNA FLJ12992 fis, clone NT2RP3000149.	2443	99
562	gi10438048	Homo sapiens	cDNA: FLJ21857 fis, clone HEP02294.	2398	96
563	gi11137965	Homo sapiens	tRNA isopentenylpyrophosphate transferase precursor RNA, complete cds.	2158	:00
563	gi7019915	Homo sapiens	cDNA FLJ20061 fis, clone COL01383.	1719	100
563	gi9803035	Caenorhabditis elegans	contains similarity to Pfam domain PF00096 (zf-C2H2), Score=12.0, E- value=1.1, N=1	407	32
564	gi7023103	Homo sapiens	cDNA FLJ10826 fis, clone NT2RP4001100.	2171	100
564	gi10434339	Homo sapiens	cDNA FLJ12690 fis, clone NT2RM4002567.	2171	100
564	gi10433458	Homo sapiens	cDNA FLJ12068 fis, clone HEMBB1002329.	2166	99
565	gi7019829	Homo sapiens	cDNA FLJ20011 fis, clone ADKA03432.	865	100
565	gi10438448	Homo sapiens	cDNA: FLJ22168 fis, clone HRC00618.	865	100
565	AAG02581	Homo sapiens	Human secreted protein, SEQ ID NO: 6662.	445	98
566	gi11558482	Homo sapiens	mRNA for B-cell lymphoma/leukaemia 11A extra long form (BCL11A-XL gene).	1543	99
566	gi12150278	Homo sapiens	C2H2-type zinc-finger protein mRNA, complete cds.	1039	99
566	gi6652688	Mus musculus	C2H2-type zinc finger protein	1033	98

100 100	SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
Homo sapiens Human secreted protein clone ydf3 1 994 100			Homo sapiens		994	
Sequence SEQ ID NO:1143. Sequence SEQ ID NO:1143. Sequence SEQ ID NO:1143. Sequence SEQ ID NO:1143. Sequence SEQ ID NO:1243. Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence SEQ ID Sequence Sequence SEQ ID S	567	AAY73435	Homo sapiens	Human secreted protein clone yd73_1	994	100
Colone DKFZp434A155), complete cds.	567	AAB43698	Homo sapiens		752	95
See	568	gi12053249	Homo sapiens	clone DKFZp434A155); complete cds.	752	95
Sequence SEQ ID NO:1143. 100	568	AAY73435		protein sequence SEQ ID NO:92.		İ
100 100	568	AAB43698	Homo sapiens	sequence SEQ ID NO:1143.		
Section	569	gi8096260	Homo sapiens	gene for Nop10p, complete cds.	344	100
	569	gi8096476	Homo sapiens	mRNA for Nop10p, complete cds.	344	100
POLR2/12 gene Polarization Pol				(H/ACA small nucleolar RNPs), clone		100
antigen protein sequence SEQ ID 578.	570	gi11595476	Homo sapiens	(POLR2J2 gene).		100
Polariza Polariza	570	AAB58870		antigen protein sequence SEQ ID 578.	409	100
SingGDS (RAP1GDS1) mRNA, alternatively spliced, complete cds. 2994 99	570	gi11595474	Homo sapiens	(POLR2J2 gene).		
571 gi13111713 Homo sapiens ,RAP1, GTP-GDP dissociation stimulator 1, clone MGC:2897, mRNA, complete cds. 2994 99 571 gi6942013 Homo sapiens exchange factor smgGDS mRNA, complete cds, alternatively spliced. 2991 99 572 gi12002978 Homo sapiens mitosin-associated protein MITAP1 (MITAP1) mRNA, complete cds. 1736 100 572 gi12043569 Homo sapiens Nudel mRNA, complete cds. 1736 100 572 gi13775593 Homo sapiens endooligopeptidase A mRNA, complete cds. 1720 99 573 gi7022325 Homo sapiens cDNA FLJ10350 fis, clone NTZRM2001131. 1243 100 573 gi3417386 Mus musculus microtubule-associated protein, MAP-115 428 48 574 gi7022502 Homo sapiens cDNA FLJ10458 fis, clone NTZRP1001457, highly similar to Homo sapiens partial mRNA for beta-transducin family protein. 2555 100 574 gi3687833 Xenopus laevis Putative Notchless protein homolog 1110 52 575 AAY51115 Homo sapiens Human HSEC6 protein.	571	gi7239381	Homo sapiens	smgGDS (RAP1GDS1) mRNA,	2995	99
Complete cds, alternatively spliced. 1736 100	571	gi13111713	Homo sapiens	, RAP1, GTP-GDP dissociation stimulator 1, clone MGC:2897, mRNA,	2994	99
MITAP1) mRNA, complete cds. 1736 100	571	gi6942013	Homo sapiens		2991	99
S72 gi13775593 Homo sapiens endooligopeptidase A mRNA, complete cds.	572	gi12002978	Homo sapiens			100
cds.	572		Homo sapiens			
NT2RM2001131.	572			cds.		
Clone DKFZp761F19121	573	gi7022325	Homo sapiens	NT2RM2001131.		<u> </u>
S74 gi7022502 Homo sapiens CDNA FLJ10458 fis, clone NT2RP1001457, highly similar to Homo sapiens partial mRNA for beta-transducin family protein. NT2RP1001457, highly similar to Homo sapiens partial mRNA for beta-transducin family protein. S74 gi12643028 Oryza sativa Putative Notchless protein homolog 1110 52	573		Homo sapiens	clone DKFZp761F19121).		<u> </u>
NT2RP1001457, highly similar to Homo sapiens partial mRNA for beta-transducin family protein. 2149 82						
Section Sect	574	gi7022502	Homo sapiens	NT2RP1001457, highly similar to Homo sapiens partial mRNA for beta-transducin	2555	100
575 AAY51115 Homo sapiens Human HSEC6 protein. 3767 99 575 gi1163174 Rattus norvegicus similar to yeast Sec6p, Swiss-Prot Accession Number P32844; similar to mammalian B94, Swiss-Prot Accession Number Q03169; Method: conceptual translation supplied by author 3606 94 575 AAB49655 Homo sapiens Human SEC7 protein sequence SEQ ID 14. 2737 89 576 gi7020303 Homo sapiens cDNA FLJ20300 fis, clone HEP06465. 1697 99	574	gi3687833		notchless	2149	82
575 gi1163174 Rattus norvegicus similar to yeast Sec6p, Swiss-Prot Accession Number P32844; similar to mammalian B94, Swiss-Prot Accession Number Q03169; Method: conceptual translation supplied by author 575 AAB49655 Homo sapiens Human SEC7 protein sequence SEQ ID 14. 576 gi7020303 Homo sapiens cDNA FLJ20300 fis, clone HEP06465. 1697 99	574	gi12643028	Oryza sativa			
norvegicus Accession Number P32844; similar to mammalian B94, Swiss-Prot Accession Number Q03169; Method: conceptual translation supplied by author Human SEC7 protein sequence SEQ ID 2737 89 14. 576 gi7C20303 Homo sapiens cDNA FLJ20300 fis, clone HEP06465. 1697 99	575	AAY51115				
575 AAB49655 Homo sapiens Human SEC7 protein sequence SEQ ID 14. 2737 89 576 gi7020303 Homo sapiens cDNA FLJ20300 fis, clone HEP06465. 1697 99		gi1163174		similar to yeast Sec6p, Swiss-Prot Accession Number P32844; similar to mammalian B94, Swiss-Prot Accession Number Q03169; Method: conceptual translation supplied by author	3606	94
	575	AAB49655	Homo sapiens		2737	89
	576	gi7020303	Homo sapiens	cDNA FLJ20300 fis, clone HEP06465.	1697	99
hydrolytic enzyme HYENZ7.		AAB67575	Homo sapiens		759	47

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
76	gi10434892	Homo sapiens	cDNA FLJ13055 fis, clone	755	47
	3 -11	•	NT2RP3001538, weakly similar to		
			HYPOTHETICAL 39.0 KD PROTEIN		
			T28D9.3 IN CHROMOSOME II.		<u> </u>
577	AAR15222	Homo sapiens	Chronic myelogenous leukaemia-derived	513	100
	ļ	-	myeloid-related protein.		
577	gi32402	Homo sapiens	Human mRNA for HP-1, a member of the	493	100
			corticostatin/defensin family.		
577	gi181527	Homo sapiens	Human neutrophil peptide (defensin) l	493	100
			mRNA, complete cds.		ļ
578	AAY41716	Homo sapiens	Human PRO860 protein sequence.	5224	100
578	AAB44272	Homo sapiens	Human PRO860 (UNQ421) protein	5224	100
			sequence SEQ ID NO:211.		
578	gi14042832	Homo sapiens	cDNA FLJ14946 fis, clone	3746	93
			PLACE2000034, weakly similar to LAR		
			PROTEIN PRECURSOR (EC 3.1.3.48).		-
579	gi7021880	Homo sapiens	cDNA FLJ10054 fis, clone	2306	100
		<u> </u>	HEMBA1001310.	2226	100
579	gi12653981	Homo sapiens	, TRIAD3 protein, clone MGC:998,	2306	100
		L	mRNA, complete cds.	2012	100
579	gi7109299	Homo sapiens	TRIAD3 mRNA, partial cds.	2013	100
580	gi3288457	Homo sapiens	mRNA for C2 domain containing PI3-	7615	99
			kinase.	2000	100
580	gi3059227	Rattus	phosphoinositide 3-kinase	3988	80
	1221222	norvegicus	Discoulation with 2 Kings C2-amount	3984	78
580	gi3041786	Mus musculus	Phosphoinositide 3-Kinase-C2gamma	1802	99
581	gi10437125	Homo sapiens	cDNA: FLJ21103 fis, clone CAS04883.	786	52
581	gi7020867	Homo sapiens	cDNA FLJ20635 fis, clone KAT03466.	297	100
582	gi13937952	Homo sapiens	, Similar to upregulated during skeletal muscle growth 5, clone MGC:14697,	291	100
			muscle growth 3, clotte MGC.14037,		
500	gi6851054	Rattus	DAPIT protein	278	91
582	g10851054	norvegicus	DAI II piotem	2.0	1
582	gi9843791	Mus musculus	stretch regulated skeletal muscle protein	259	84
583	gi7582286	Homo sapiens	BM-007	599	100
583	AAG02907	Homo sapiens	Human secreted protein, SEQ ID NO:	477	98
383	AAG02907	Homo sapiens	6988.] 7,7	1
583	gi3878572	Caenorhabditis	M01F1.6	161	28
203	g13676372 .	elegans	14101111.0	10,	20
584	gi13477103	Homo sapiens	, clone MGC:1012, mRNA, complete cds.	3001	99
584	gi12052999	Homo sapiens	mRNA; cDNA DKFZp434E1711 (from	2619	98
J0 4	g112032333	Tiomo suprens	clone DKFZp434E1711); complete cds.		1,0
584	gi7020996	Homo sapiens	cDNA FLJ20721 fis, clone HEP15722.	2402	100
585	AAW48892	Homo sapiens	Human guanylate binding protein B	2645	94
363	77.4440072	Home suprems	(HGBPB).	20.0	
585	gi12803663	Homo sapiens	, guanylate binding protein 1, interferon-	2000	66
202	g112003003	Trome suprem	inducible, 67kD, clone MGC:3949,		1
			mRNA, complete cds.		
585	gi183002	Homo sapiens	Human guanylate binding protein isoform	2000	66
505	51105002	Trome suprem	I (GBP-2) mRNA, complete cds.		
586 gi	gi7023366	Homo sapiens	cDNA FLJ10983 fis, clone	3218	99
	g., -25500	110.111	PLACE1001781, weakly similar to		
]		PROBABLE	1	
			PHOSPHOMANNOMUTASE (EC	1	
			5.4.2.8).		
586	gi12052930	Homo sapiens	mRNA; cDNA DKFZp566B1524 (from	3216	99
555	5-1202730		clone DKFZp566B1524); complete cds.	1	1
586	gi3395586	Schizosacchar	similarity to phosphomannomutases	1211	43
	1 2	omyces pombe		1	

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
587	gi13537208	Mus musculus	Mel18 and Bmil like ring finger	347	40
587	gi2440074	Homo sapiens	mRNA for RNF3A (DONG1) ring finger protein.	347	37
587	gi13537206	Homo sapiens	hMBLR mRNA, complete cds.	345	40
588	gi14042249	Homo sapiens	cDNA FLJ14610 fis, clone NT2RP1000958, weakly similar to AUTOANTIGEN NGP-1.	2797	99
588	gi14042246	Homo sapiens	cDNA FLJ14608 fis, clone NT2RP1000915, weakly similar to AUTOANTIGEN NGP-1.	2741	99
588	gi6457340	Homo sapiens	E2IG3 (E2IG3) mRNA, complete cds.	2650	100
589	gi7020925	Homo sapiens	cDNA FLJ20673 fis, clone KAIA4464.	2232	100
589	gi7682684	Homo sapiens	phosphoprotein associated with GEMs (PAG) mRNA, complete cds.	2222	99
589	gi7707799	Rattus norvegicus	Csk binding protein Cbp	1696	78
590	gi6682873	Homo sapiens	rec mRNA, complete cds.	2002	100
590	gi7230612	Rattus norvegicus	small rec	1916	95
590	gi3881771	Caenorhabditis elegans	contains similarity to Pfam domain: PF01529 (DHHC zinc finger domain), Score=137.4, E-value=8.4e-38, N=1	586	39
591	gi439522	Mus musculus	ribosomal protein S3	678	100
591	gi57728	Rattus rattus	ribosomal protein S3 (AA 1-243)	678	100
591	gi13111933	Homo sapiens	, ribosomal protein S3, clone MGC:3657, mRNA, complete cds.	678	100
592	gi6599070	Homo sapiens	mRNA for LIM domains containing protein 1.	3675	99
592	gi6599307	Mus musculus	LIM domains containing protein 1	2728	76
592	gi13548632	Homo sapiens	partial LIMD1 gene for LIM domains containing 1, exons 1-2, complete sequence.	2690	99
593	gi7020974	Homo sapiens	cDNA FLJ20706 fis, clone KAIA1273.	2824	98
593	gi12082725	Mus musculus	B cell phosphoinositide 3-kinase adaptor	411	29
593	AAG02945	Homo sapiens	Human secreted protein, SEQ ID NO: 7026.	526	100
594	gi11596144	Homo sapiens	STE20-like kinase mRNA, partial cds.	5159	99
594	gi3452473	Rattus norvegicus	scrinc/threonine protein kinase TAO1	5117	98
594	AAY55937	Homo sapiens	Human SULU3 protein.	4045	100
595	gi695802	Homo sapiens	transcription factor SL1 mRNA, partial cds.	1693	99
595	gi1842206	Mus musculus	TAFI68	1326	76
596	gi7020363	Homo sapiens	cDNA FLJ20335 fis, clone HEP11429.	2940	99
596	AAB65680	Homo sapiens	Novel protein kinase, SEQ ID NO: 208.	2940	99
596	AAB32078	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 136.	826	100
597	gi7020747	Homo sapiens	cDNA FLJ20558 fis, clone KAT11870.	2990	100
597	gi12053175	Homo sapiens	mRNA; cDNA DKFZp434A172 (from clone DKFZp434A172); complete cds.	2990	100
597	gi10439123	Homo sapiens	cDNA: FLJ22650 fis, clone HSI07344.	2166	100
598	gi7023601	Homo sapiens	cDNA FLJ11127 fis, clone PLACE1006225.	1897	100
598	gi12224968	Homo sapiens	mRNA; cDNA DKFZp667E105 (from clone DKFZp667E105).	620	100
598	gi14043433	Homo sapiens	, clone IMAGE:3952677, mRNA, partial cds.	549	41
599	gi6483296	Homo sapiens	CDH9 mRNA for cadherin-9, complete cds.	4132	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
599	gi867999	Gallus gallus	chicken cadherin-6B	3044	72
599	gi974185	Homo sapiens	mRNA for cadherin-6, complete cds.	3032	72
600	gi5734605	Homo sapiens	mRNA for KARP-1-binding protein 3, complete cds.	750	51
600	gi5734601	Homo sapiens	mRNA for KARP-1-binding protein 1 (KAB1), complete cds.	750	51
600	gi5734603	Homo sapiens	mRNA for KARP-1-binding protein 2 (KAB2), complete cds.	750	51
601	gi10434848	Homo sapiens	cDNA FLJ13028 fis, clone NT2RP3001055, weakly similar to Drosophila melanogaster separation anxiety protein (san) mRNA.	889	100
601	gi10435107	Homo sapiens	cDNA FLJ13194 fis, clone NT2RP3004378, weakly similar to Drosophila melanogaster separation anxiety protein (san) mRNA.	889	100
601	AAB56739	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1317.	874	98
602	gi13325182	Homo sapiens	, clone IMAGE:3638994, mRNA, partial cds.	897	100
602	gi12654203	Homo sapiens	, clone IMAGE:3449323, mRNA, partial cds.	560	100
602	gi4514314	Bacillus halodurans	YlqF	260	39
603	gi10954046	Homo sapiens	oxidation protection protein (OXR1) mRNA, complete cds.	1034	97
603	gi13540300	Mus musculus	nucleolar protein C7B	1431	94
603	gi7021988	Homo sapiens	cDNA FLJ10125 fis, clone HEMBA1002954.	1441	99
604	gil150495	Mus musculus	homology to nucleosome assembly proteins; specifically expressed in neurons	211	36
604	gi1161252	Glycine max	nucleosome assembly protein 1	136	40
604	gi5931610	Homo sapiens	mRNA for Nucleosome Assembly Protein 1-like 2, complete cds.	196	37
605	gi7547029	Homo sapiens	GAP-like protein (N61) mRNA, complete cds.	4684	99
605	gi7688683	Homo sapiens	kinesin heavy chain-like protein (KHCHP) mRNA, complete cds.	822	100
605	AAG03378	Homo sapiens	Human secreted protein, SEQ ID NO: 7459.	633	99
606	gi7022593	Homo sapiens	cDNA FLJ10511 fis, clone NT2RP2000656.	1425	100
606	gi12224996	Homo sapiens	mRNA; cDNA DKFZp667G248 (from clone DKFZp667G248).	1031	100
606	gi10436327	Homo sapiens	cDNA FLJ13991 fis, clone Y79AA1002115.	803	100
607	gi8885998	Rattus norvegicus	neuronal C-SRC tyrosine-specific protein kinase	2826	98
607	gi201057	Mus musculus	tyrosine-specific protein kinase	2822	98
607	gi338460	Homo sapiens	Human c-src-1 proto-oncogene, exon 12.	2815	98
608	gi7243633	Homo sapiens	RB-associated KRAB repressor (RBAK) mRNA, complete cds.	3993	100
608	gi7243635	Mus musculus	RB-associated KRAB repressor	3025	78
608	gi10434235	Homo sapiens	cDNA FLJ12629 fis, clone NT2RM4001828, moderately similar to	1881	73
(00	-:7000:00		ZINC FINGER PROTEIN 84.	082	100
609	gi7008402	Homo sapiens	kappa B-ras 1 mRNA, complete cds.	982	100
609	gi14042659	Homo sapiens	cDNA FLJ14843 fis, clone	978	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			PLACE1000040, weakly similar to TRANSFORMING PROTEIN P21/K- RAS 2B.		
	gi7239257	Mus musculus	kappaB-Rasi	952	94
609	gi13625164	Homo sapiens	ankyrin mRNA, complete cds.	426	100
610	gi13623164 gi12698638	Homo sapiens	ankyrin-repeat family A protein 2 (ANKRA2) mRNA, complete cds.	426	100
610	gi10434525	Homo sapiens	CDNA FLJ12814 fis, clone NT2RP2002520, weakly similar to Homo sapiens transcription factor RFX-B (RFXB) mRNA.	426	100
611	gi7959841	Homo sapiens	PRO1853	510	100
611	AAG01282	Homo sapiens	Human secreted protein, SEQ ID NO: 5363.	301	100
612	gi5757703	Mus musculus	syntrophin-associated serine-threonine protein kinase	7464	92
612	gi13537204	Homo sapiens	mRNA for MAST205, complete cds.	4616	68
612	gi406058	Mus musculus	protein kinase	4569	65
613	gi7020724	Homo sapiens	cDNA FLJ20545 fis, clone KAT11476.	1780	100
613	AAB63186	Homo sapiens	Human secreted protein sequence encoded by gene 3 SEQ ID NO:112.	1693	100
613	gi7243701	Drosophila melanogaster	WDS	1574	91
614	gi13383476	Homo sapiens	NUB1 (NUB1) mRNA, complete cds.	3109	100
614	gi5360093	Homo sapiens	NY-REN-18 antigen mRNA, complete cds.	2958	95
614	gi863014	Mus musculus	BS4 peptide	2671	84
615	AAB87345	Homo sapiens	Human gene 4 encoded secreted protein HDPFY41, SEQ ID NO:86.	4534	100
615	gi4886489	Homo sapiens	mRNA; cDNA DKFZp564L2123 (from clone DKFZp564L2123); partial cds.	2892	99
615	gi12711793	Homo sapiens	estrogen regulated LIV-1 protein (LIV-1) mRNA, complete cds.	1171	39
616	gi7638247	Homo sapiens	mesenchymal stem cell protein DSCD75 mRNA, complete cds.	1063	100
616	gi12654929	Homo sapiens	, mesenchymal stem cell protein DSCD75, clone MGC:5515, mRNA, complete cds.	1063	100
616	AAB03956	Homo sapiens	Human mesenchymal stem cell polypeptide.	1063	100
617	gi7582304	Homo sapiens	BM-016	584	100
617	AAW78199	Homo sapiens	Human secreted protein encoded by gene 74 clone HGBAC11.	562	98
617	AAW85610	Homo sapiens	Secreted protein clone eh80_1.	562	98
618	gi13603398	Homo sapiens	mRNA for SEZ6L, complete cds.	4199	98
618	gi13185723	Homo sapiens	n 1755 can be A, G, C, or T	2164	49
618	AAB70537	Homo sapiens	Human PRO7 protein sequence SEQ ID NO:14.	2164	49
619	gi3880445	Caenorhabditis elegans	contains similarity to Pfam domain: PF02214 (K+ channel tetramerisation domain), Score=79.5, E-value=2.3e-20, N=1	327	40
619	AAY34129	Homo sapiens	Human potassium channel K+Hnov28.	195	40
619	AAZ11907_	Homo sapiens	Human potassium channel K+Hnov28 cDNA (5' splice variant 1).	195	40
620	gi10437116	Homo sapiens	cDNA: FLJ21097 fis, clone CAS03931.	1146	100
620	gi14250732	Homo sapiens	, chromosome 11 open reading frame 14, clone MGC:12847, mRNA, complete cds.		100
620	gi13276621	Homo sapiens	mRNA; cDNA DKFZp761G1913 (from	378	43

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
1.01			clone DKFZp761G1913).		
621	gi10437078	Homo sapiens	cDNA: FLJ21069 fis, clone CAS01594.	955	58
621	gi5911935	Homo sapiens	mRNA; cDNA DKFZp586N1922 (from clone DKFZp586N1922); partial cds.	867	100
621	AAB27870	Homo sapiens	Protein fragment encoded by gene 27.	657	100
622	gi13097159	Homo sapiens	, tumor protein, translationally-controlled 1, clone MGC:5308, mRNA, complete cds.	898	100
622	gi14043771	Homo sapiens	, clone MGC:14243, mRNA, complete cds.	898	100
622	gi7573519	Homo sapiens	TPT1 gene for translationally controlled tumor protein (TCTP), exons 1-6.	898	100
623	gi7020339 ·	Homo sapiens	cDNA FLJ20320 fis, clone HEP08923.	1135	100
623	AAB18972	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1135	100
623	gi1314162	Schizosacchar omyces pombe	seven transmembrane protein	217	29
624	gi6467990	Mus musculus	PDZ domain actin binding protein Shroom	4816	66
624	gi6467992	Mus musculus	actin binding protein ShroomS	4816	66
624	gi13938323	Homo sapiens	, Similar to shroom, clone IMAGE:3349317, mRNA, partial cds.	4006	99
625	gi12804029	Homo sapiens	, clone IMAGE:3940519, mRNA, partial cds.	1551	100
625	AAY21850	Homo sapiens	Human signal peptide-contianing protein (SIGP) (clone ID 1880830).	1109	100
625	gi8655657	Homo sapiens	mRNA; cDNA DKFZp762O076 (from clone DKFZp762O076).	593	57
626	gi7328140	Homo sapiens	mRNA; cDNA DKFZp762D096 (from clone DKFZp762D096); partial cds.	601	100
626	gi13436341	Homo sapiens	, Similar to RIKEN cDNA 1600014C10 gene, clone MGC:10922, mRNA, complete cds.	384	100
627	gi1293559	Mus musculus	astrotactin	4312	95
627	gi6502571	Mus musculus	astrotactin2	2580	51
627	gi6502573	Homo sapiens	astrotactin2 (ASTN2) mRNA, complete cds.	2569	51
628	AAY73387	Homo sapiens	HTRM clone 3340290 protein sequence.	1439	95
628	AAY48312	Homo sapiens	Human prostate cancer-associated protein 9.	1073	84
628	gi12654077	Homo sapiens	, clone IMAGE:3458173, mRNA, partial cds.	1045	86
629	gi11095188	Homo sapiens	dipeptidyl peptidase 8 (DPP8) mRNA, complete cds.	3521	99
629	gi14042790	Homo sapiens	cDNA FLJ14920 fis, clone PLACE1007416, weakly similar to DIPEPTIDYL PEPTIDASE IV (EC 3.4.14.5).	2457	99
629	gi7020273	Homo sapiens	cDNA FLJ20283 fis, clone HEP04088.	2483	100
630	gil1095188	Homo sapiens	dipeptidyl peptidase 8 (DPP8) mRNA, complete cds.	2560	99
630	gi14042790	Homo sapiens	cDNA FLJ14920 fis, clone PLACE1007416, weakly similar to DIPEPTIDYL PEPTIDASE IV (EC 3.4.14.5).	2457	99
630	gi11095192	Homo sapiens	dipeptidyl peptidase 8 (DPP8) mRNA, partial cds, alternatively spliced.	2482	100
631	gi7020611	Homo sapiens	cDNA FLJ20481 fis, clone KAT07534.	2211	99
631	AAY57908	Homo sapiens	Human transmembrane protein HTMPN-	975	44

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			32.		
631	AAB54284	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:736.	516	40
632	gi35700	Homo sapiens	Human mRNA for phosphoriobosyl pyrophosphate synthetase subunit II (EC 2.7.6.1).	1596	99
632	gi206434	Rattus norvegicus	phosphoribosyl pyrophosphate synthetase II	1585	98
632	gi56979	Rattus norvegicus	ribose-phosphate pyrophosphokinase subunit II (AA 1-318)	1585	98
(22	gi11181620	Homo sapiens	Rag D mRNA, complete cds.	1276	100
633 633	gi6808148	Homo sapiens	mRNA; cDNA DKFZp761H171 (from clone DKFZp761H171); partial cds.	1276	100
633	AAB56443	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1021.	1276	100
634	gi6807893	Homo sapiens	mRNA; cDNA DKFZp434H2226 (from clone DKFZp434H2226); partial cds.	1079	100
635	gi10435042	Homo sapiens	cDNA FLJ13152 fis, clone NT2RP3003385, highly similar to Mus musculus SKD3 mRNA.	3495	100
635	gi4958935	Rattus norvegicus	suppressor of potassium transport defect 3	3085	88
635	gi563129	Mus musculus	SKD3	3066	88
636	AAB20322	Homo sapiens	Human protein phosphatase and kinase protein-1.	1770	100
636	gi1903458	Dictyostelium discoideum	myosin heavy chain kinase B	236	32
636	gi2104701	Mus musculus	elongation factor-2 kinase	199	29
637	gi7670003	Homo sapiens	mRNA; cDNA DKFZp434P0531 (from clone DKFZp434P0531).	1850	100
637	gi7417474	Homo sapiens	chromosome 14 clone RP11-493G17 and CTD-2516D11 map 14q24.3, complete sequence.	1251 .	49
637	gi7018538	Homo sapiens	mRNA; cDNA DKFZp434P0111 (from clone DKFZp434P0111); partial cds.	330	43
638	gi7022367	Homo sapiens	cDNA FLJ10375 fis, clone NT2RM2001950.	3056	100
638	AAY53026	Homo sapiens	Human secreted protein clone cn922_5 protein sequence SEQ ID NO:58.	1752	95
638	gi4336692	Drosophila melanogaster	Abnormal X segregation	816	37
639	gi7020972	Homo sapiens	cDNA FLJ20705 fis, clone KAIA1571.	3641	99
639	gi12007334	Homo sapiens	IRS-1 PH domain binding protein PHIP mRNA, complete cds.	3632	99
639	gi14286226	Homo sapiens	, pleckstrin homology domain interacting protein, clone MGC:15187, mRNA, complete cds.	3632	99
640	gi7689025	Homo sapiens	uncharacterized hypothalamus protein HT013 mRNA, complete cds.	978	96
641	gi9937505	Homo sapiens	PLIC-2 mRNA, complete cds.	3167	100
641	gi6563288	Homo sapiens		3162	99
641	AAB47122	Homo sapiens		3162	99
641	AAY53001	Homo sapiens	1 004 1	811	100
642	AAG01114	Homo sapiens		641	99
642	gi 12652989	Homo sapiens	2011	489	57
643	gi7021064	Homo sapiens		2240	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
543	gi10438264	Homo sapiens	cDNA: FLJ22019 fis, clone HEP07982.	2187	98
643	gi577428	Rattus norvegicus	Ca2+-dependent activator protein; calcium-dependent actin-binding protein	1787	77
644	gi7023651	Homo sapiens	cDNA FLJ11159 fis, clone PLACE1006966.	2865	99
644	gi7023118	Homo sapiens	cDNA FLJ10835 fis, clone NT2RP4001210.	1253	100
644	gi600058	Saccharomyce s cerevisiae	N1342	710	39
645	gi7020012	Homo sapiens	cDNA FLJ20121 fis, clone COL05942.	1334	99
646	gi14336697	Homo sapiens	16p13.3 sequence section 2 of 8.	609	100
646	gi13436122	Homo sapiens	, non-metastatic cells 4, protein expressed in, clone MGC:11088, mRNA, complete cds.	609	100
646	gi1945762	Homo sapiens	H.sapiens mRNA for nucleoside- diphosphate kinase.	609	100
647	AAB24225	Homo sapiens	Human vesicle associated protein 4 SEQ ID NO:4.	2946	99
647	gi10439139	Homo sapiens	cDNA: FLJ22662 fis, clone HSI08080.	2703	99
647	AAB58427	Homo sapiens	Lung cancer associated polypeptide sequence SEQ ID 765.	1711	99
648	gi7020604	Homo sapiens	cDNA FLJ20477 fis, clone KAT07271.	2639	99
648	gi6672090	Drosophila melanogaster	Vegetable	578	32
649	gi12802986	Homo sapiens	, ring finger protein 24, clone MGC:1815, mRNA, complete cds.	811	100
649	gi5420200	Homo sapiens	Novel human mRNA from chromosome 20, similar to SW:GOLI_DROME Q06003 GOLIATH PROTEIN.	811	100
649	gi5102892	Homo sapiens	mRNA full length insert cDNA clone EUROIMAGE 566628.	566	100
650	gi6841346	Homo sapiens	HSPC054	497	98
651	gi7209305	Homo sapiens	mRNA for FLJ00002 protein, partial cds.	7637	100
651	gi6599226	Homo sapiens	mRNA; cDNA DKFZp434L0827 (from clone DKFZp434L0827); partial cds.	3519	100
651	gi10440406	Homo sapiens	mRNA for FLJ00036 protein, partial cds.	3457	99
652	gi7018505	Homo sapiens	mRNA; cDNA DKFZp434E2220 (from clone DKFZp434E2220).	2470	100
652	gi14042579	Homo sapiens	cDNA FLJ14796 fis, clone NT2RP4001235.	2466	99
652	gi7018507	Homo sapiens	mRNA; cDNA DKFZp434O0420 (from clone DKFZp434O0420).	2466	99
653	gi552196	Plasmodium lophurae	histidine-rich protein	192	40
653	gi160362	Plasmodium falciparum	knob protein	178	42
653	gi3845095	Plasmodium falciparum	knob-associated His-rich protein	172	40
654	AAY70539	Homo sapiens	Human Factor 8 Homologue.	1353	83
654	gi14043498	Homo sapiens	, Similar to neuropilin 1, clone MGC:12920, mRNA, complete cds.	189	34
654	gi7271465	Homo sapiens	soluble neuropilin-1 mRNA, complete cds.	189	34
655	gi7019959	Homo sapiens	cDNA FLJ20087 fis, clone COL03793.	3964	100
655	gi13569705	Homo sapiens	channel kinase 2 (CHAK2) mRNA, complete cds.	3942	99
655	AAY95433	Homo sapiens	Human calcium channel SOC-2/CRAC-1 C-terminal polypeptide.	1172	71
656	gi6094668	Homo sapiens	BAC clone RP11-343N14 from 2,	208	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			complete sequence.		
656	gi10435833	Homo sapiens	cDNA FLJ13729 fis, clone PLACE3000121, weakly similar to VESICULAR TRAFFIC CONTROL PROTEIN SEC15.	208	100
656	gi2827162	Rattus norvegicus	rsec15	160	73
657	gi10434153	Homo sapiens	cDNA FLJ12580 fis, clone NT2RM4001116, weakly similar to HYPOTHETICAL 216.3 KD PROTEIN R06F6.8 IN CHROMOSOME II.	1806	99
657	gi12053255	Homo sapiens	mRNA; cDNA DKFZp434D105 (from clone DKFZp434D105); complete cds.	1806	99
657	gi5901808	Drosophila melanogaster	BcDNA.GH03694	619	56
658	gi11181618	Homo sapiens	Rag C mRNA, complete cds.	2072	100
658	gi12007486	Homo sapiens	GTPase-interacting protein 2 mRNA, complete cds.	2069	99
658	gi13529335	Mus musculus	Similar to Rag C protein	2039	98
659	gi13537208	Mus musculus	Mel18 and Bmil like ring finger	347	40
659	gi2440074	Homo sapiens	mRNA for RNF3A (DONG1) ring finger protein.	347	37
659	gi13537206	Homo sapiens	hMBLR mRNA, complete cds.	345	40
660	gi7023690	Homo sapiens	cDNA FLJ11184 fis, clone PLACE1007507.	1043	99
661	gi7020878	Homo sapiens	cDNA FLJ20641 fis, clone KAT02782.	2552	99
661	gi11992034	Rattus norvegicus	antisense RNA overlapping MCH protein	1609	65
662	AAB56646	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1224.	915	98
662	gi12053357	Homo sapiens	mRNA; cDNA DKFZp586G2122 (from clone DKFZp586G2122); complete cds.	900	100
662	AAB36598	Homo sapiens	Human FLEXHT-20 protein sequence SEQ ID NO:20.	791	59
663	AAW93947	Homo sapiens	Human regulatory molecule HRM-3 protein.	1732	100
663	gi3288459	Homo sapiens	mRNA for transcription elongation factor TFIIS.h.	1673	100
663	gi3288547	Mus musculus	transcription elongation factor TFIIS.h	1543	90
664	gi14042893	Homo sapiens	cDNA FLJ14984 fis, clone Y79AA1000349, highly similar to M.musculus Spnr mRNA for RNA binding protein.	3478	100
664	gi13377630	Homo sapiens	spermatid perinuclear RNA-binding protein mRNA, complete cds.	3459	99
664	gi12053237	Homo sapiens	mRNA; cDNA DKFZp434N214 (from clone DKFZp434N214); complete cds.	3406	100
665	gi10436573	Homo sapiens	cDNA FLJ14183 fis, clone NT2RP2004920, weakly similar to TRANSCRIPTIONAL REGULATOR ATRX.	4423	99
665	gi10434345	Homo sapiens	cDNA FLJ12693 fis, clone NT2RP1000324.	4369	99
665	AAB27235	Homo sapiens	Human EXMAD-13 SEQ ID NO: 13.	3346	100
666	gi9858154	Homo sapiens	tubby super-family protein (TUSP) mRNA, complete cds, alternatively spliced.	3598	100
666	gi9502082	Homo sapiens	tubby super-family protein (TUSP) mRNA, complete cds.	3556	100

SEQ ID	Accession No.	Species	Description	Score	% Identity
NO:	gi9502080	Mus musculus	tubby super-family protein	3505	98
566	gi7106796	Horno sapiens	HSPC203	554	100
67	gi9963859	Homo sapiens	PTD019 mRNA, complete cds.	554	100
567 567	AAY35987	Homo sapiens	Extended human secreted protein sequence, SEQ ID NO. 236.	554	100
	<u> </u>			3398	99
668 668	gi6996442 gi6996589	Homo sapiens Rattus	CTL1 gene. CTL1 protein	3291	96
668	gi6996587	norvegicus Torpedo	CTL1 protein	2454	71
669	gi6808165	Homo sapiens	mRNA; cDNA DKFZp761A052 (from clone DKFZp761A052).	2265	100
	-:10430059	Homo sapiens	cDNA: FLJ22607 fis, clone HSI04846.	1992	100
669	gi10439058	Mus musculus	DXImx46e protein	1958	98
669 670	gi7673616 gi6808252	Homo sapiens	mRNA; cDNA DKFZp434D1319 (from clone DKFZp434D1319); partial cds.	2336	100
		-	N-75	221	27
670	gi170035	Glycine max	pre-pro polypeptide (AA -25 to 284)	219	27
670 671	gi18576 AAW93947	Glycine max Homo sapiens	Human regulatory molecule HRM-3	1116	99
671	gi3288459	Homo sapiens	protein. mRNA for transcription elongation factor TFIIS h.	1057	99
671	gi3288547	Mus musculus	transcription elongation factor TFIIS.h	950	86
672	gi10434615	Homo sapiens	cDNA FLJ12875 fis, clone NT2RP2003777.	1818	99
672	gi8778741	Arabidopsis thaliana	T30E16.12	254	27
672	gi6520214	Arabidopsis thaliana	ZCF61	228	29
673	AAB88424	Homo sapiens	Human membrane or secretory protein clone PSEC0197.	3032	99
673	gi9294464	Arabidopsis thaliana	long-chain-fatty-acid-CoA ligase-like	581	37
673	gi699196	Mycobacteriu m leprae	4-coumarate-coA ligase	326	45
674	gi7022969	Homo sapiens	cDNA FLJ10747 fis, clone NT2RP3001799.	3378	99
674	AAY86211	Homo sapiens	Nuclear transport protein clone hfb066 protein sequence.	1432	87
674	gi10439560	Homo sapiens	cDNA: FLJ23007 fis, clone LNG00451.	703	100
674 675	gi7021968	Homo sapiens	cDNA FLJ10111 fis, clone HEMBA1002696.	2753	99
	114017762	Mus musculus	FLJ10111	2214	92
675	gi14017768		cDNA: FLJ23501 fis, clone LNG02837.	2160	90
675 676	gi10440211 gi7021968	Homo sapiens Homo sapiens	cDNA FLJ10111 fis, clone HEMBA1002696.	2728	98
	1	1 due manufact	FLJ10111	2200	90
676	gi14017768	Mus musculus	cDNA: FLJ23501 fis, clone LNG02837.	2237	92
676	gi10440211	Homo sapiens	cDNA FLJ20036 fis, clone COL00219.	2834	100
677 677	gi7019869 gi12723779	Homo sapiens Lactococcus	UNKNOWN PROTEIN	306	35
		lactis subsp. lactis		7007	20
677	gi8885520	Streptococcus gordonii	streptococcal hemagglutinin	297	29
678	gi10437508	Homo sapiens	cDNA: FLJ21415 fis, clone COL04030.	1129	100
679	gi3135314	Homo sapiens	chromosome 7q22 sequence, complete sequence.	1226	100
679	gi6752287	Homo sapiens	Novel human gene mapping to chomosome X.	390	43

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
679	AAB28327	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 111.	265	100
680	gi3135314	Homo sapiens	chromosome 7q22 sequence, complete sequence.	1199	95
680	gi6752287	Homo sapiens	Novel human gene mapping to chomosome X.	363	41
680	AAB28327	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 111.	265	100
681	gi10439089	Homo sapiens	cDNA: FLJ22626 fis, clone HSI06109.	2120	99
681	gi11044557	Homo sapiens	Human DNA sequence from clone RP11- 42415 on chromosome 6 Contains a novel gene, STSs, GSSs and a CpG island, complete sequence.	1185	64
681	gi12654241	Homo sapiens	, Similar to splicing factor, arginine/serine-rich 4 (SRp75), clone MGC:5283, mRNA, complete cds.	949	98
682	gi14042277	Homo sapiens	cDNA FLJ14626 fis, clone NT2RP2000288.	3029	99
682	gi7022410	Homo sapiens	cDNA FLJ10402 fis, clone NT2RM4000457.	2279	100
682	gi6841196	Homo sapiens	HSPC273	1086	100
683	gi2815604	Homo sapiens	Opa-interacting protein OIP2 mRNA, partial cds.	1364	100
683	AAB63276	Homo sapiens	Human breast cancer associated antigen protein sequence SEQ ID NO:638.	839	96
683	AAB63406	Homo sapiens	Human breast cancer associated antigen protein sequence SEQ ID NO:768.	839	96
684	AAB07228	Homo sapiens	Human prostate cancer predisposing protein HPC2.	4325	100
684	AAY99850	Homo sapiens	Human sulphatase G.	4315	99
684	gi10946497	Pan troglodytes	ELAC2	4283	98
685	gi7688979	Homo sapiens	uncharacterized bone marrow protein BM042	895	100
685 .	AAB36580	Homo sapiens	Human FLEXHT-2 protein sequence SEQ ID NO:2.	895	100
685	AAB34771	Homo sapiens	Human secreted protein fragment encoded by DNA clone vq23 1.	888	99
686	gi10438990	Homo sapiens	cDNA: FLJ22559 fis, clone HSI01591.	1897	100
686	gi8954034	Arabidopsis thaliana	F10K1.17	162	31
687	gi7020674	Homo sapiens	cDNA FLJ20515 fis, clone KAT09889.	2027	100
687	AAB20331	Homo sapiens	Human protein phosphatase and kinase protein-10.	1472	92
687	AAB73226	Homo sapiens	Human phosphatase NP 060746 h.	576	83
688	gi6688145	Homo sapiens	mRNA for NICE-3 protein, clone 1023j12.	1019	100
688	gi4689120	Homo sapiens	HSPC012	717	93
688	gi12655055	Homo sapiens	, DKFZP586G1722 protein, clone MGC:1147, mRNA, complete cds.	717	93
689	gi7023701	Homo sapiens	cDNA FLJ11190 fis, clone PLACE1007583.	1317	100
690	gi6469703	Mycobacteriu m tuberculosis	DipZ	203	31
691	gi13676779	Mus musculus	Arkadia	1939	93
691	gi13752369	Gallus gallus	ring finger protein	1888	91
691	gi13752371	Xenopus laevis	ring finger-H2 protein	1537	76
692	gi458255	Homo sapiens	Human X-linked PEST-containing	2849	99

SEQ ID	Accession No.	Species	Description	Score	% Identity
NO:	110.		transporter (XPCT) gene, exon 6.		
592	gi458247	Homo sapiens	Human X-linked PEST-containing transporter (XPCT) mRNA, partial cds.	2766	99
592	gi2944356	Mus musculus	X-linked PEST-containing transporter	2249	88
693	gi14042736	Homo sapiens	cDNA FLJ14888 fis, clone PLACE1003762.	2034	99
593	gi6841178	Homo sapiens	HSPC264	2019	99
694	gi7023413	Homo sapiens	cDNA FLJ11012 fis, clone PLACE1003190, weakly similar to SOF1 PROTEIN.	2377	99
694	gi14042745	Homo sapiens	cDNA FLJ14893 fis, clone PLACE1004302, weakly similar to SOF1 PROTEIN.	2377	99
694	gi5912184	Homo sapiens	mRNA; cDNA DKFZp564O0463 (from clone DKFZp564O0463); partial cds.	1159	99
695	gi7022931	Homo sapiens	cDNA FLJ10724 fis, clone NT2RP3001176.	2683	99
695	gi14198202	Mus musculus	Similar to melanoma antigen recognized by T cells 2	2126	82
695	gi4826524	Homo sapiens	Novel human gene mapping to chomosome 1.	982	92
696	gi7022990	Homo sapiens	cDNA FLJ10761 fis, clone NT2RP3004669, weakly similar to ETHANOLAMINE KINASE (EC 2.7.1.82).	2119	99
696	gi9998952	Homo sapiens	ethanolamine kinase (EKII) mRNA, complete cds.	930	56
696	gi532128	Drosophila melanogaster	ethanolamine kinase	525	45
697	gi186774	Homo sapiens	Human Kruppel related zinc finger protein (HTF10) mRNA, complete cds.	986	38
697	gi5441615	Canis familiaris	zinc finger protein	988	37
697	gi38032	Homo sapiens	Human ZNF43 mRNA.	947	36
698	gi13537202	Homo sapiens	PC-LKC mRNA for protocadherin LKC, complete cds.	2877	100
698 .	gi7020017	Homo sapiens	cDNA FLJ20124 fis, clone COL06056.	2862	99
698	AAY01410	Homo sapiens	Secreted protein encoded by gene 28 clone HE9ND43.	963	100
699	gi7688977	Homo sapiens	uncharacterized bone marrow protein BM041	888	100
699	AAY86515	Homo sapiens	Human gene 71-encoded protein fragment, SEQ ID NO:430.	888	100
699	gi7018421	Homo sapiens	mRNA; cDNA DKFZp564J157 (from clone DKFZp564J157).	880	99
700	gi7209307	Homo sapiens	mRNA for FLJ00003 protein, partial cds.	1102	100
700	gi14276857	Homo sapiens	PC2-glutamine-rich-associated protein (PCQAP) mRNA, complete cds.	429	93
700	gi14043091	Homo sapiens	, clone IMAGE:3350171, mRNA, partial cds.	429	93
701	gi7020678	Homo sapiens	cDNA FLJ20517 fis, clone KAT10235.	2821	99
701	gi10177966	Arabidopsis thaliana	uridine kinase-like protein	1068	44
701	gi496728	Saccharomyce s cerevisiae		775	37
702	gi7022789	Homo sapiens	cDNA FLJ10634 fis, clone NT2RP2005654, weakly similar to CYSTEINE STRING PROTEIN.	1512	100
702	AAB67446	Homo sapiens	Amino acid sequence of a human	1512	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			chaperone polypeptide.		ļ
702	AAG01952	Homo sapiens	Human secreted protein, SEQ ID NO: 6033.	422	98
703	gi7021321	Homo sapiens	Gemin4 mRNA, complete cds.	5481	99
703	gi10945430	Homo sapiens	chromosome 17 clone PAC P579 HC90, HC71AC, HC6 and HC56 genes, complete sequence.	5452	100
703	gi7018412	Homo sapiens	mRNA; cDNA DKFZp434D174 (from clone DKFZp434D174).	4359	99
704	gi9964287	Homo sapiens	hypertension-related calcium-regulated gene mRNA, complete cds.	1129	100
704	gi10434820	Homo sapiens	cDNA FLJ13008 fis, clone NT2RP3000456.	1129	100
704	gi12803673	Homo sapiens	, HT002 protein; hypertension-related calcium-regulated gene, clone MGC:3418, mRNA, complete cds.	1129	100
705	gi10435947	Homo sapiens	cDNA FLJ13814 fis, clone THYRO1000368.	3588	99
705	gi3878402	Caenorhabditis elegans	similar to C2 domain	300	25
705	gi3002479	Leishmania major	L3162.1	198	25
706	gi11907998	Homo sapiens	BCL-6 corepressor (BCOR) mRNA, complete cds; alternatively spliced.	2449	100
706	gi7020277	Homo sapiens	cDNA FLJ20285 fis, clone HEP04260.	1131	99
706	gi10432606	Homo sapiens	cDNA FLJ11362 fis, clone HEMBA1000244.	458	50
707	gi7768662	Homo sapiens	C4ST mRNA for chondroitin 4-sulfotranseferase, complete cds.	1870	100
707	gi8925966	Homo sapiens	chondroitin 4-O-sulfotransferase 1 mRNA, complete cds.	1870	100
707	gi7572958	Homo sapiens	mRNA for chondroitin-4-sulfotransferase (C4ST gene).	1865	99
708	gi2731561	Homo sapiens	ATP receptor subunit (P2X5) mRNA, complete cds.	2167	96
708	gi1552522	Homo sapiens	Human ionotropic ATP receptor P2X5a mRNA, complete cds.	2131	96
708	gi3387944	Homo sapiens	clone 24793 ionotropic ATP receptor P2X5b mRNA, complete cds.	1608	99
709	gi7021105	Homo sapiens	cDNA FLJ20793 fis, clone COL00343.	1587 435	100
709	gi7206854	Caenorhabditis elegans	contains similarity to Pfam family PF00085 (Thioredoxins), Score 113, E=9.6e-33, N=1		
709	gi13775331	Caenorhabditis elegans	contains similarity to Pfam family PF00085 (Thioredoxin), score=320.7, E=1.8e-95, N=3	297	28
710	AAY04315	Homo sapiens	Human secreted protein encoded by gene 23.	385	100
710	AAB12155	Homo sapiens	Hydrophobic domain protein isolated from HT-1080 cells.	385	100
711	gi13624098	Homo sapiens	cervical cancer 1 protooncogene protein p40 mRNA, complete cds.	520	100
711	gi12653253	Homo sapiens	, DKFZP586A011 protein, clone MGC:8483, mRNA, complete cds.	520	100
711	gi4886473	Homo sapiens	mRNA; cDNA DKFZp586A011 (from clone DKFZp586A011); partial cds.	520	100
712	gi927415	Homo sapiens	H. sapiens mRNA for carnitine acetyltransferase.	3209	98
712	gi13879380	Mus musculus	Similar to carnitine acetyltransferase	3010	90

1713 gi738632 Mus musculus carntine acetyltransferase 2967 89 89713 19437507 Homo sapiens TERA 1198 100 100 101 10	SEQ ID	Accession No.	Species	Description	Score	% Identity
1713 194379507 Homo sapiens TERA 1198 100 1713 1910439906 Homo sapiens CDNA: FLIZ3279 fis, clone HEP06870 1198 100 1714 1910439906 Homo sapiens CDNA: FLIZ3279 fis, clone MGC:1093, 1198 100 1714 1911498104 Homo sapiens CDNA: FLIZ3279 fis, clone MGC:1093, 1198 100 1714 1911498104 Homo sapiens CDNA: FLIZ10964 fis, clone 1196 100 1714 1911498104 Homo sapiens CDNA: FLIZ10969 fis, clone 661 100 1715 1715 171020019 Homo sapiens CDNA: FLIZ11269 fis, clone 661 100 1715 171020019 Homo sapiens CDNA: FLIZ10125 fis, clone 2740 100 1715 171020019 Homo sapiens CDNA: FLIZ10125 fis, clone 2740 100 1715 171020019 Homo sapiens GDNA: FLIZ10125 fis, clone 2740 100 1715 1716 171020019 Homo sapiens GDNA: FLIZ10125 fis, clone 2740 100 1716 1717 1718 171			Mus musculus	carnitine acetyltransferase	2967	
1713					1198	100
198						
196 100				, TERA protein, clone MGC:1093,		
Title	714	gi7023336	Homo sapiens	cDNA FLJ10964 fis, clone	1196	100
Till	714	gi14198104	Homo sapiens	, clone MGC:16981, mRNA, complete	1196	100
AAB67579	714	gi7023823	Homo sapiens	cDNA FLJ11269 fis, clone	661	100
Polymorphic Polymorphic	715	AAB67579	Homo sapiens	hydrolytic enzyme HYENZ11.	<u> </u>	
715	715	gi7020019	Homo sapiens	cDNA FLJ20125 fis, clone COL06152.		
716		gi13527857		pol polyprotein	298	26
716 AAW53863 Homo sapiens Human gravin polypeptide. 8868 99 716 AAB15380 Homo sapiens Human gravin protein sequence. 8868 99 717 gi7021891 Homo sapiens Human gravin protein sequence. 2306 99 717 gi10432215 Homo sapiens CDNA FLJ10606 fis, clone 1959 86 717 gi14042890 Homo sapiens CDNA FLJ14982 fis, clone 1959 86 718 gi6224691 Homo sapiens CDNA FLJ14982 fis, clone 1959 86 718 AAB36158 Homo sapiens Novel human transporter SUT-1 (SUT-1) 3271 100 718 AAB23625 Homo sapiens Novel human transporter protein SEQ ID 3268 99 719 gi7020123 Homo sapiens Human secreted protein SEQ ID NO: 50. 3268 99 719 gi34328904 Homo sapiens Human fGIF. 1262 99 719 gi4522409 Homo sapiens Human FGIF. 1262 99 720 </td <td>716</td> <td>gi2218077</td> <td></td> <td>gravin mRNA, complete cds.</td> <td></td> <td></td>	716	gi2218077		gravin mRNA, complete cds.		
Tile		AAW53863	Homo sapiens			
Tit				Human gravin protein sequence.		
HEMBA1006789. Hemba 1006789. Hemba1006789. Hemba1006789. Hemba1006789. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Hemba1000258. Na+i/sulfate cotransporter SUT-1 (SUT-1) 3271 100 mRNA, complete cds. Novel human transporter protein SEQ ID 3268 99 Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human transporter protein SEQ ID Novel human SEQ ID Novel human transporter protein SEQ ID Novel human	717	gi7021891	Homo sapiens	HEMBA1001407.		99
Y79AA1000258. Y79AA1000258	717	-		HEMBA1006789.		
mRNA, complete cds.	717	·		Y79AA1000258.		
NO: 2.	718	gi6224691		mRNA, complete cds.		
719 gi7020123 Homo sapiens cDNA FLJ20189 fis, clone COLF0657. 1264 99 719 gi14328904 Homo sapiens fetal globin-:nducing factor (FGIF) mRNA, complete cds. 1262 99 719 AAB71861 Homo sapiens Human FGIF. 1262 99 720 gi6690250 Homo sapiens clone HQ0659 PRO0659 mRNA, complete cds. 926 100 720 gi12654109 Homo sapiens , PRO0659 protein, clone MGC:4888, mRNA, complete cds. 926 100 721 gi608025 Homo sapiens Luman ankyrin G (ANK-3) mRNA, complete cds. 580 32 721 gi3885972 Rattus norvegicus 270 kDa ankyrin G isoform 575 32 721 gi178646 Homo sapiens Homan erythroid ankyrin mRNA, complete cds. 609 35 722 gi7020915 Homo sapiens CDNA FLJ20666 fis, clone KAIA608. 1229 96 722 gi3169096 Schizosacchar omyces pombe possible pre-mRNA processing by similarity to yeast prp39 420 37 723 gi10434720	718			NO: 2.		
719 gi14328904 Homo sapiens fetal globin-:nducing factor (FGIF) mRNA, complete cds. 1262 99 719 AAB71861 Homo sapiens Human FGIF. 1262 99 720 gi6690250 Homo sapiens clone HQ0659 PRO0659 mRNA, complete cds. 926 100 720 gi12654109 Homo sapiens ,PRO0659 protein, clone MGC:4888, mRNA, complete cds. 926 100 721 gi608025 Homo sapiens Human ankyrin G (ANK-3) mRNA, complete cds. 580 32 721 gi3885972 Rattus norvegicus 270 kDa ankyrin G isoform 575 32 721 gi178646 Homo sapiens Human erythroid ankyrin mRNA, complete cds. 609 35 722 gi7020915 Homo sapiens cDNA FLJ20666 fis, clone KAIA608. 1229 96 722 gi1458279 Caenorhabditis elegans contains sim:larity to TPR domains 252 29 723 gi7020729 Homo sapiens cDNA FLJ120548 fis, clone KAT11542. 2200 100 723 gi10434720 Homo sapiens	718			Human secreted protein SEQ ID NO: 50.		
mRNA, complete cds. 1262 99						
T20 gi6690250 Homo sapiens Clone HQ0659 PRO0659 mRNA, complete cds.	719			mRNA, complete cds.		
Complete cds. Complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MGC:4888, mRNA, complete cds. PRO0659 protein, clone MSNA, complete cds. PRO0659 protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grade protein grad	719					
mRNA, complete cds. 1280 32 32 32 32 32 32 32 3	720			complete cds.		
Complete cds. 270 kDa ankyrin G isoform 575 32	720			mRNA, complete cds.	i	
1721 gi178646 Homo sapiens Human erythroid ankyrin mRNA, 609 35 1722 gi7020915 Homo sapiens cDNA FLJ20666 fis, clone KAIA608. 1229 96 1723 gi1458279 Caenorhabditis elegans cDNA FLJ20548 fis, clone KAT11542. 2200 100 1724 gi1967781 Homo sapiens ANKRD2 gene for skeletal muscle ankyrin repeat, exons 1-9. 2903 99 1724 gi10433458 Homo sapiens cDNA FLJ12068 fis, clone 2903 99 1726 Gi10433458 Homo sapiens cDNA FLJ12068 fis, clone 2903 99 1727 Gi10433458 Homo sapiens cDNA FLJ12068 fis, clone 2903 99 1728 Gi10433458 Homo sapiens cDNA FLJ12068 fis, clone 2903 99 1729 Gi10433458 Homo sapiens cDNA FLJ12068 fis, clone 2903 99	721		•	complete cds.		
Complete cds. Complete cds			norvegicus			
Total Process Proces	721			complete cds.		
Omyces pombe Similarity to yeast prp39						
Proceed	722		omyces pombe	similarity to yeast prp39		
T23 gi10434720 Homo sapiens CDNA FLJ12942 fis, clone NT2RP2005139, weakly similar to 2-5A-DEPENDENT RIBONUCLEASE (EC 3.1.26). T23 gi11967781 Homo sapiens ANKRD2 gene for skeletal muscle ankyrin repeat, exons 1-9. T24 gi10433458 Homo sapiens CDNA FLJ12068 fis, clone LEMBB1002329. P9	722	l	elegans			
NT2RP2005139, weakly similar to 2-5A-DEPENDENT RIBONUCLEASE (EC 3.1.26).	723	gi7020729				
723 gi11967781 Homo sapiens ANKRD2 gene for skeletal muscle ankyrin repeat, exons 1-9. 174 30 724 gi10433458 Homo sapiens cDNA FLJ12068 fis, clone HEMBB1002329. 2903 99	723	gi10434720	Homo sapiens	NT2RP2005139, weakly similar to 2-5A- DEPENDENT RIBONUCLEASE (EC 3.1.26).	2200	100
724 gi10433458 Homo sapiens cDNA FLJ12068 fis, clone HEMBB1002329. 2903 99	723	gi11967781	Homo sapiens	ANKRD2 gene for skeletal muscle	174	30
	724	gi10433458	Homo sapiens	cDNA FLJ12068 fis, clone	2903	99
1/24 g110434339 Homo sapiens CDNA FLI 1/2090 IIS, Clone 2898 99	724	gi10434339	Homo sapiens	cDNA FLJ12690 fis, clone	2898	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
110.			NT2RM4002567.		
724	gi10436665	Homo sapiens	cDNA FLJ14252 fis, clone OVARC1001341.	2167	99
725	gi10434638	Homo sapiens	cDNA FI J12889 fis, clone NT2RP2004098, weakly similar to ADENYLATE CYCLASE (EC 4.6.1.1).	3026	100
725	gi14250313	Homo sapiens	, clone MGC:16864, mRNA, complete cds.	3026	100
725	gi7020356	Homo sapiens	cDNA FLJ20331 fis, clone HEP10410.	1914	99
726	AAY13947	Homo sapiens	Human transmembrane protein, HP10495.	655	100
726	AAY07878	Homo sapiens	Human secreted protein fragment encoded from gene 27.	655	100
726	gi6841296	Homo sapiens	HSPC323	449	85
727	gi7159733	Homo sapiens	mRNA for ETAA16 protein.	4318	100
727	AAB10622	Homo sapiens	Human Ewing tumor associated antigen protein.	4318	100
728	gi7020138	Homo sapiens	cDNA FLJ20199 fis, clone COLF1162.	2123	99
728	AAY91948	Homo sapiens	Human cytoskeleton associated protein 3 (CYSKP-3).	1650	99
728	gi7020210	Homo sapiens	cDNA FLJ20246 fis, clone COLF6458.	1474	99
729	gi13182775	Homo sapiens	CDA11 mRNA, complete cds.	1495	99
729	gi13937914	Homo sapiens	, clone MGC:12519, mRNA, complete cds.	973	97
729	gi2257524	Schizosacchar omyces pombe	HYPOTHETICAL 47.4KD PROTEIN IN SHP1-SEC17 INTERGENIC REGION	536	42
730	gi7020242	Homo sapiens	cDNA FLJ20265 fis, clone COLF9334.	2813	99
730	gi14042159	Homo sapiens	cDNA FLJ14559 fis, clone NT2RM2001998.	2812	99
730	gi499005	Saccharomyce s cerevisiae	HRC830	128	32
731	gi7022375	Homo sapiens	cDNA FLJ10379 fis, clone NT2RM2002014.	3182	99
731	gi14010930	Homo sapiens	BAC clone RP11-576F1 from 2, complete sequence.	1868	100
731	gi1573555	Haemophilus influenzae Rd	transcription accessory protein (tex)	691	42
732	gi10434409	Homo sapiens	cDNA FLJ12737 fis, clone NT2RP2000337.	1001	99
733	gi7019597	Homo sapiens	clone PAC 270M7 chromosome 21 map 21q11.2, complete sequence.	5944	100
733	gi7407669	Homo sapiens	chromosome 21 PAC 30P13 map 21q11.2, complete sequence, containing gene for nuclear factor RIP140.	5944	100
733	gi7717256	Homo sapiens	chromosome 21 segment HS21C007.	5944	100
734	gi7021956	Homo sapiens	cDNA FLJ10103 fis, clone HEMBA1002495, weakly similar to LIGHT-MEDIATED DEVELOPMENT PROTEIN DET1.	1415	100
734	AAB64828	Homo sapiens	Human secreted protein sequence encoded by gene 12 SEQ ID NO:114.	869	99
734	gi4038594	Lycopersicon esculentum	tDET1 protein	413	37
735	gi6752405	Streptococcus pneumoniae	PspA	137	24
736	gi5080758	Homo sapiens	chromosome 19, BAC 331191 (CIT-B-471f3), complete sequence.	1486	55
736	gi456269	Mus musculus domesticus	zinc finger protein 30	1478	54

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
736	gi4567179	Homo sapiens	chromosome 19, BAC 37295 (CIT-B-21A4), complete sequence.	1281	62
737	gi7023220	Homo sapiens	cDNA FL/10893 fis, clone NT/2RP4002791.	4557	99
737	gi14042072	Homo sapiens	cDNA FLJ14507 fis, clone NT2RM1000399.	4439	97
737	gi7582296	Homo sapiens	BM-012	1807	99
738	gil 1596985	Homo sapiens	chromosome 14 clone RP11-361H10 map 14q24.3, complete sequence.	1751	100
738	gi7020945	Homo sapiens	cDNA FLJ20689 fis, clone KAIA2890.	1738	99
738	gi6067151	Homo sapiens	chromosome 14 BAC 98L12, complete sequence.	1159	100
739	gi6941888	Homo sapiens	ubiquitin-specific processing protease (USP25) mRNA, complete cds.	5638	99
739	AAB31550	Homo sapiens	A human ubiquitin specific protease (USP).	5638	99
739	gi6693824	Homo sapiens	ubiquitin-specific protease (USP21) mRNA, complete cds.	4022	99
740	gi6693824	Homo sapiens	ubiquitin-specific protease (USP21) mRNA, complete cds.	5465	99
740	AAB31546	Homo sapiens	A human ubiquitin specific protease 25 (USP25).	5465	99
740	AAF24881_ aal	Homo sapiens	DNA encoding a human ubiquitin specific protease 25 (USP25).	5465	99
741	gi7161175	Homo sapiens	mRNA for 19A24 protein (19A24 gene).	1726	100
741	gi13021810	Homo sapiens	NK cell receptor (CS1) mRNA, complete cds.	1349	100
741	AAB32373	Homo sapiens	Human secreted protein sequence encoded by gene 3 SEQ ID NO:59.	1349	100
742	gi7023747	Homo sapiens	cDNA FLJ11219 fis, clone PLACE1008122.	2553	100
742	gi7022222	Homo sapiens	cDNA FLJ10287 fis, clone HEMBB1001387.	880	97
742	AAG01392	Homo sapiens	Human secreted protein, SEQ ID NO: 5473.	569	99
743	gi7023747	Homo sapiens	cDNA FLJ11219 fis, clone PLACE1008122.	2442	97
743	gi7022222	Homo sapiens	cDNA FLJ10287 fis, clone HEMBB1001387.	769	89
743	AAG01392	Homo sapiens	Human secreted protein, SEQ ID NO: 5473.	569	99
744	gi6434857	Homo sapiens	pallid mRNA, complete cds.	872	100
744	gi13435969	Homo sapiens	, pallid (mouse) homolog, pallidin, clone MGC:4983, mRNA, complete cds.	872	100
744	gi6456870	Mus musculus	syntaxin 13-interacting protein pallid	754	87
745	gi6841480	Homo sapiens	HSPC129	2378	99
745	gi6841354	Homo sapiens	HSPC058	1825	99
745	gi7022613	Homo sapiens	cDNA FLJ10523 fis, clone NT2RP2000863.	1489	
746	gi7023644	Homo sapiens	cDNA FLJ11155 fis, clone PLACE1006935.	1826	99
746	AAB18981	Homo sapiens	Amino acid sequence of a human transmembrane protein.	1000	99
746	gi13384531	Caenorhabditis elegans	similar to C. elegans protein T16H12.10	680	40
747	gi13544089	Homo sapiens	, clone IMAGE:4053618, mRNA, partial cds.	2749	99
747	gi6007859	Chlamydomon as reinhardtii	dynein heavy chain alpha	246	30

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
747	gi2065436	Schizosacchar omyces pombe	tealp	227	28
748	gi6650778	Homo sapiens	PRO1575	297	100
	gi8926849	Homo sapiens	mRNA for Pex3p, complete cds.	1892	99
749	gi4092648	Homo sapiens	mRNA for PEX3 protein, partial.	1892	99
749 749	gi4092048 gi4218426	Homo sapiens	pex3 gene (joined CDS, promoter and exon 1).	1892	99
	1200000	Mus musculus	early B-cell factor	3064	99
750 750	gi309209 gi6630994	Homo sapiens	early B-cell transcription factor (EBF)	3033	98
			mRNA, partial cds.	3023	97
750	gi7687988	Gallus gallus	early B-cell factor	3102	99
751	gi10436636	Homo sapiens	cDNA FLJ14228 fis, clone NT2RP3004148.		
751	gi14278861	Homo sapiens	PHD zinc finger transcription factor mRNA, complete cds.	2127	100
751	gi12804495	Homo sapiens	, clone IMAGE:3356959, mRNA, partial cds.	1472	100
752	gi6594639	Homo sapiens	dynein intermediate chain DNAII (DNAII) mRNA, complete cds.	1773	100
752	gi6635422	Homo sapiens	dynein intermediate chain DNAII	1768	99
132	g10033422	Tromo sapiens	(DNAI1) gene, exon 20 and complete cds.		
752	gi927637	Anthocidaris crassispina	dynein intermediate chain 2	961	61
753	gi5924385	Rattus norvegicus	ribosomal protein S271	412	100
753	gi12803647	Homo sapiens	, ribosomal protein S27 (metallopanstimulin 1), clone MGC:3659, mRNA, complete cds.	412	100
753	gi1373421	Homo sapiens	Human ribosomal protein S27 mRNA, complete cds.	412	100
754	gi1655432	Mus musculus	plexin 2	9646	96
754 754	gi6010215	Homo sapiens	mRNA for partial OCT/plexin-A2 protein.	6985	99
754	gi1665757	Mus musculus	plexin 1	6359	63
754	gi7770189	Homo sapiens	PRO2325	901	100
755 756	gi77022885	Homo sapiens	cDNA FLJ10697 fis, clone NT2RP3000527, weakly similar to ZINC FINGER PROTEIN 43.	3318	99
756	gi10434872	Homo sapiens	cDNA FLJ13043 fis, clone NT2RP3001338, weakly similar to ZINC FINGER PROTEIN 81.	957	43
756	gi38032	Homo sapiens	Human ZNF43 mRNA.	346	25
756 757	gi14042238	Homo sapiens	cDNA FLJ14604 fis, clone NT2RP1000363, moderately similar to R.norvegicus LL5 mRNA.	1107	93
757	AAB43723	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1168.	647	86
757	gi14044043	Homo sapiens	, clone IMAGE:4299555, mRNA, partial cds.	467	66
766	gi7106766	Homo sapiens	HSPC188	532	100
758			, clone MGC:4355, mRNA, complete cds.	529	99
758 758	gi12804349 gi1002516	Saccharomyce		115	27
759	gi6175593	s cerevisiae Homo sapiens	transcription factor IIIC90 mRNA,	4326	99
760	gi7023345	Homo sapiens	complete cds. cDNA FLJ10970 fis, clone	647	99
	AAG03409	Homo sapiens	PLACE1000948. Human secreted protein, SEQ ID NO:	239	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			7490.		
761	gi5441541	Canis familiaris	Ribosomal protein	447	94
761	gi304526	Cricetulus griseus	ribosomal protein S17	447	94
761	gi10439453	Homo sapiens	cDNA: FLJ22917 fis, clone KAT06430.	447	94
762	gi6635353	Homo sapiens	RU1 (RU1) mRNA, complete cds.	4638	99
762	gi8100079	Mus musculus	polycomb-group proteins	4176	88
762	gi8100077	Rattus norvegicus	polycomb-group protein	4152	88
763	gi12804681	Homo sapiens	, \$100 calcium-binding protein, beta (neural), clone MGC:1323, mRNA, complete cds.	479	100
763	gi337730	Homo sapiens	Human S100 protein beta-subunit gene, exon 3.	479	100
763	gi404769	Mus musculus	S100 beta protein	473	98
764	gi7106782	Homo sapiens	HSPC196	617	98
764	gi7106786	Homo sapiens	HSPC198	617	98
764	AAW74871	Homo sapiens	Human secreted protein encoded by gene 143 clone HBMDM46.	617	98
765	gi3851206	Homo sapiens	chromosome 19, cosmid F19847, complete sequence.	1282	100
765	gi13276629	Homo sapiens	mRNA; cDNA DKFZp761D221 (from clone DKFZp761D221); complete cds.	815	35
765	gi5701573	Caenorhabditis elegans	similar to S. pombe phosphoprotein (GB:X86179)	430	33
766	gi7020238	Homo sapiens	cDNA FLJ20262 fis, clone COLF7748.	1393	100
766	gi12653607	Homo sapiens	, clone IMAGE:3162218, mRNA, partial cds.		98
766	AAY86358	Homo sapiens	Human gene 11-encoded protein fragment, SEQ ID NO:273.	996	95
767	gi2588619	Homo sapiens	BAC clone CTB-104F4 from 7q21-q22, complete sequence.	2037	100
767	gi1707507	Homo sapiens	H.sapiens mRNA for mitochondrial transcription termination factor.	2037	100
767	gi12654289	Homo sapiens	, transcription termination factor, mitochondrial, clone MGC:5000, mRNA, complete cds.	2033	99
768	gi1314373	Homo sapiens	Human aquaporin-5 (AQP5) gene, exon 4 and complete cds.	1336	100
768	gi664760	Rattus norvegicus	aquaporin-5	1245	91
768	gi4894460	Mus musculus	aquaporin 5	1235	91
769	gi13097624	Homo sapiens	, clone IMAGE:3608084, mRNA, partial cds.	1093	100
769	gi10438279	Homo sapiens	cDNA: FLJ22029 fis, clone HEP08661.	615	60
769	gi13325154	Homo sapiens	, clone IMAGE:3635709, mRNA, partial cds.	609	45
770	AAB48789	Homo sapiens	Human prostate cancer-predisposing protein, CA7 CG04.	2878	100
770	gi11321424	Mus musculus	Ral-A exchange factor RalGPS2	2073	96
770	gi7637906	Homo sapiens	Ral guanine nucleotide exchange factor RalGPS1A mRNA, complete cds.	1224	70
771	gi13623239	Homo sapiens	, Similar to SGC32445 protein, clone MGC:10610, mRNA, complete cds.	1080	99
771	gi7547035	Homo sapiens	SGC32445 protein (SGC32445) mRNA, complete cds.		100
771	gi10434977	Homo sapiens	cDNA FLJ13110 fis, clone NT2RP3002549, moderately similar to	519	64

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			HYPOTHETICAL 26.6 KD PROTEIN T19C3.4 IN CHROMOSOME III.		
172	gi13939858	Homo sapiens	RITA	2614	100
72	gi10048470	Homo sapiens	C2H2-like zinc finger protein (ZNF463) mRNA, complete cds.	2614	100
772	gi8575775	Homo sapiens	KRAB zinc finger protein (RITA) mRNA, complete cds.	2614	100
773	gi12654989	Homo sapiens	, clone MGC:5623, mRNA, complete cds.	2300	100
773	gi3329425	Homo sapiens	huntingtin interacting protein HYPE mRNA, partial cds.	963	100
773	gi429189	Haemophilus somnus	surface protein	152	41
774	gi14028017	Mesorhizobiu m loti	argininosuccinate lyase	199	26
774	gi2182606	Rhizobium sp. NGR234	Y4rH	179	29
775	gi3098311	Oryctolagus cuniculus	elongation factor 1 A2	2410	100
775	gi8886507	Homo sapiens	elongation factor 1 A-2 (EF1A-2) gene, complete cds.	2410	100
775	gi12653327	Homo sapiens	, eukaryotic translation elongation factor 1 alpha 2, clone MGC:8362, mRNA, complete cds.	2410	100
776	gi6624095	Homo sapiens	BAC clone RP11-294L11 from 2, complete sequence.	2515	97
776	AAY66674	Homo sapiens	Membrane-bound protein PRO1277.	2515	97
776	AAB87542	Homo sapiens	Human PRO1277.	2515	97
777	gi6049162	Homo sapiens	rhabdoid tumor deletion region protein 1 (RTDR1) mRNA, complete cds.	1732	100
777	gi14290442	Homo sapiens	, rhabdoid tumor deletion region protein 1, clone MGC:16968, mRNA, complete cds.	1732	100
778	AAB66071	Homo sapiens	Human INTERCEPT 296	1787	99
778	AAB18992	Homo sapiens	Amino acid sequence of a human transmembrane protein.	880	58
778	AAB26325	Homo sapiens	Human CASB618 protein.	880	58
779	gi643656	Rattus norvegicus	synaptotagmin VII	1851	95
779	gi12667446	Rattus norvegicus	synaptotagmin VIIs	1851	95
779	gi6136786	Mus musculus	synaptotagmin VII	1842	95
780	gi7020988	Homo sapiens	cDNA FLJ20716 fis, clone HEP19742.	1048	100
780	gi4033606	Adiantum capillus-veneris	Extensin	131	38
780	gi169347	Phaseolus vulgaris	hydroxyproline-rich glycoprotein	130	38
781	gi7020477	Homo sapiens	cDNA FLJ20401 fis, clone KAT00901	1644	96
781	gi7022002	Homo sapiens	cDNA FLJ10135 fis, clone HEMBA1003117.	590	40
781	gi7022284	Homo sapiens	cDNA FLJ10324 fis, clone NT2RM2000567.	590	40
782	gi6808186	Homo sapiens	mRNA; cDNA DKFZp434D0218 (from clone DKFZp434D0218), partial cds.	1322	99
783	gi505544	Homo sapiens	H.sapiens mRNA for Zinc-finger protein (ZNFpT1).	1211	99
783	AAY58627		Protein regulating gene expression 688 PRGE-20.		50
783	gi9187356	Homo sapiens	mRNA full length insert cDNA clone	687	50

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			EUROIMAGE 2107395.		<u> </u>
784	gi8895094	Homo sapiens	SH3-containing protein SH3GLB2 mRNA, complete cds.	1975	97
784	gi4929591	Homo sapiens	CGI-61 protein mRNA, complete cds.	706	69
784	gi8896092	Homo sapiens	SH3-containing protein SH3GLB1 mRNA, complete cds.	706	69
785	gi7770175	Homo sapiens	PRO2249	1827	99
785	gi11527602	Homo sapiens	mRNA for MCM10 homolog, complete cds.	1827	99
785	gi12053187	Homo sapiens	mRNA, cDNA DKFZp434H152 (from clone DKFZp434H152); complete cds.	1682	99
786	gi7023364	Homo sapiens	cDNA FLJ10982 fis, clone PLACE1001692, moderately similar to S- ACYL FATTY ACID SYNTHASE THIOESTERASE, MEDIUM CHAIN (EC 3.1.2.14).	1413	99
786	gi7023563	Homo sapiens	cDNA FLJ11106 fis, clone PLACE1005763, moderately similar to S- ACYL FATTY ACID SYNTHASE THIOESTERASE, MEDIUM CHAIN (EC 3.1.2.14).	1099	98
786	gi205326	Rattus norvegicus	S-acyl fatty acid sunthetase thio ester hydrolase, medium chain	807	55
787	gi2599502	Homo sapiens	protocadherin 68 (PCH68) mRNA, complete cds.	327	43
787	AAY24913	Homo sapiens	Human ontherin.	327	43
787	AAY94991	Homo sapiens	Human secreted protein vc35_1, SEQ ID NO:22.	296	28
788	gi7023688	Homo sapiens	cDNA FLJ11183 fis, clone PLACE1007488, weakly similar to PUTATIVE RHO/RAC GUANINE NUCLEOTIDE EXCHANGE FACTOR.	2260	100
788	gi3342246	Rattus norvegicus	actin-filament binding protein Frabin	725	32
788	gi595425	Homo sapiens	Human faciogenital dysplasia (FGD1) mRNA, complete cds.	759	32
789	gi6554165	Homo sapiens	receptor protein tyrosine phosphatase (RPTP-rho) mRNA, alternatively spliced, complete cds.	7734	99
789	gi13378306	Mus musculus	brain RPTPmam4 isoform I	7499	97
789	gi32456	Homo sapiens	H.sapiens hR-PTPu gene for protein tyrosine phosphatase.	4995	64
790	gi7020479	Homo sapiens	cDNA FLJ20402 fis, clone KAT00919.	2024	99
790	gi7770205	Homo sapiens	PRO2521	1957	97
790	gi10241843	Mus musculus	gasdermin	282	29
791	gi5262472	Homo sapiens	mRNA; cDNA DKFZp564J102 (from clone DKFZp564J102); partial cds.	1602	100
792	gi10436457	Homo sapiens	cDNA FLJ14084 fis, clone HEMBB1002383.	830	100
792	AAY94940	Homo sapiens	Human secreted protein clone yi62_1 protein sequence SEQ ID NO:86.	830	100
792	AAY57922	Homo sapiens	Human transmembrane protein HTMPN-46.	830	100
793	gi7328061	Homo sapiens	mRNA; cDNA DKFZp76112312 (from clone DKFZp76112312); partial cds.	2723	100
793	gi14039825	Mus musculus	gamma-1 syntrophin	2579	93
793	gi8247279	Homo sapiens	mRNA for syntrophin 4.	2271	97
794	g:6164674	Homo sapiens	heterogeneous nuclear ribonucleoprotein, alternate transcript (RALY) mRNA,	730	66

SEQ ID	Accession	Species	Description	Score	% Td===4:4
NO:	No.	 		 	Identity
794	gi14250048	Homo sapiens	complete cds. , heterogeneous nuclear ribonucleoprotein C (C1/C2), clone MGC:14574, mRNA, complete cds.	705	53
794	gi13937888	Homo sapiens	, Similar to heterogeneous nuclear ribonucleoprotein C, clone MGC:12469, mRNA, complete cds.	704	53
795	gi12653905	Homo sapiens	, Similar to Max dimerization protein 3, clone MGC:2383, mRNA, complete cds.	1045	100
795	AAY93137	Homo sapiens	Human Myx protein.	1023	98
795	AAB35713	Homo sapiens	Human Mad3 protein sequence.	1010	97
796	gi7020704	Homo sapiens	cDNA FLJ20533 fis, clone KAT10931.	585	98
797	gi7106878	Homo sapiens	HSPC244	398	98
797	AAY07855	Homo sapiens	Human secreted protein fragment encoded from gene 4.	398	98
797	gi13274582	Mus musculus	thymus atrophy-related protein	383	95
798	gi8886483	Gallus gallus	EURL	1178	74
798	gi10435877	Homo sapiens	cDNA FLJ13763 fis, clone PLACE4000089.	873	98
798	AAG01108	Homo sapiens	Human secreted protein, SEQ ID NO: 5189.	561	100
799	AAY33297	Homo sapiens	Human membrane spanning protein MSP-4.	.781	100
799	AAB61149	Homo sapiens	Human NOV18 protein.	781	100
799	AAB61150	Homo sapiens	Human NOV19 protein.	781	100
800	gi8099348	Homo sapiens	zinc finger protein (ZFP) mRNA, 4066 complete cds.		99
800	gi2293535	Homo sapiens	zinc finger protein (ZnF20) mRNA, complete cds.	1863	49
800	gi11527849	Mus musculus	zinc finger protein SKAT2	1323	58
801	gi7023523	Homo sapiens	cDNA FLJ11082 fis, clone PLACE1005206.	2693	99
801	gi9558010	Leishmania major	possible cDNA flj11082 fis, clone place1005206	134	26
802	gi6841558	Homo sapiens	HSPC168	1502	100
802	gi6453346	Homo sapiens	Novel human gene on chromosome 20.	1502	100
802	gi13542748	Mus musculus	RIKEN cDNA 3230401D17 gene	1314	86
803	gi7020468	Homo sapiens	cDNA FLJ20396 fis, clone KAT00561.	931	100
803	AAB18980	Homo sapiens	Amino acid sequence of a human transmembrane protein.	931	100
803	AAY91632	Homo sapiens	Human secreted protein sequence encoded by gene 25 SEQ ID NO:305.	914	98
804	gi6650345	Homo sapiens	alpha-catenin-like protein VR22 mRNA, complete cds.	4478	99
804	gi222788	Gallus gallus	alpha N-catenin	2765	60
804	AAR58778	Homo sapiens	Neural alpha-catenin protein.	2765	60
805	gi10434911	Homo sapiens			38
805	gi5912258	Homo sapiens	mRNA; cDNA DKFZp586K0524 (from clone DKFZp586K0524); partial cds.	190	41
805	gi7022673	Homo sapiens	cDNA FLJ10562 fis, clone NT2RP2002701	154	44
806	gi10435877	Homo sapiens	cDNA FLJ13763 fis, clone PLACE4000089.	876	99
806	gi8886483	Gallus gallus	EURL	868	72
806	AAG01108	Homo sapiens	Human secreted protein, SEQ ID NO:	561	100
	J		5189.		L

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
807	gi4521254	Mus musculus	cornichon-like protein	867	100
807	AAB60464	Homo sapiens	Human cell cycle and proliferation protein CCYPR-12, SEQ ID NO:12.	729	81
807	AAY76218	Homo sapiens	Human secreted protein encoded by gene 95.	716	81
808	gi7407144	Homo sapiens	protocadherin Fat 2 (FAT2) mRNA, complete cds.	22667	99
808	gi3449286	Rattus norvegicus	MEGF1	18806	81
808	gi6688786	Mus musculus	mouse fat 1 cadherin	8928	47
809	gi7407144	Homo sapiens	protocadherin Fat 2 (FAT2) mRNA, complete cds.	19770	99
809	gi3449286	Rattus norvegicus	MEGF1	16567	82
809	gi6688786	Mus musculus	mouse fat 1 cadherin	8928	47
810	gi7020201	Homo sapiens	cDNA FLJ20241 fis, clone COLF6335.	2420	100
810	gi10435321	Homo sapiens	cDNA FLJ13337 fis, clone OVARC1001880.	1279	99
810	gi7020600	Homo sapiens	cDNA FLJ20475 fis, clone KAT07206.	634	60
811	gi6483290	Homo sapiens	CDH7 mRNA for cadherin-7, complete cds.	4032	100
811	gi10803408	Homo sapiens	mRNA for cadherin-7 (CDH7 gene).	3965	98
811	gi868001	Gallus gallus	chicken cadherin-7	3830	93
812	gi13276621	Homo sapiens	mRNA; cDNA DKFZp761G1913 (from clone DKFZp761G1913).	1204	97
812	gi8977983	Mus musculus	neuronal interacting factor X 1 (NIX1)	699	78
812	gi10437116	Homo sapiens	cDNA: FLJ21097 fis, clone CAS03931.	297	42
814	gi13279269	Homo sapiens	, clone IMAGE:3631943, mRNA, partial cds.	1480	100
814	gi6808028	Homo sapiens	mRNA; cDNA DKFZp761C029 (from clone DKFZp761C029); partial cds.	857	100
814	AAW88657	Homo sapiens	Secreted protein encoded by gene 124 clone HPMCJ92.	436	94
815	gi7959853	Homo sapiens	PRO1966	281	100
816	gi7259234	Mus musculus	contains transmembrane (TM) region	718	65
816	AAY94954	Homo sapiens	Human secreted protein clone iw66_1 protein sequence SEQ ID NO:114.	679	58
816	AAB62810	Homo sapiens	Human nervous system associated protein NSPRT3 amino acid sequence.	678	61
817	gi5921144	Schizosacchar omyces pombe	mip1	1489	48
817	gi458938	Saccharomyce s cerevisiae	Yhr186cp	469	30
817	gi9366720	Trypanosoma brucei	possible t16o11.22 protein.	277	45
819	gi7020799	Homo sapiens	cDNA FLJ20590 fis, clone KAT09052.	727	100
820	gi7020555	Homo sapiens	cDNA FLJ20449 fis, clone KAT05575.	1857	99
820	AAY79269	Homo sapiens	Human testis-specific transcription factor PHELIX.	1696	99
821	gi6482350	Homo sapiens	CAC-1 mRNA, partial cds.	1136	100
821	gi13937595	Homo sapiens	, Similar to RIKEN cDNA 1810017F10 gene, clone MGC:2583, mRNA, complete	560	94
821	AAY25770	Homo sapiens	cds. Human secreted protein encoded from gene 60.	560	94
822	gi10434608	Homo sapiens	gene 60. cDNA FLJ12871 fis, clone 2023 NT2RP2003751.		100
822	gi6093227	Homo sapiens	mRNA; cDNA DKFZp434I0850 (from clone DKFZp434I0850); partial cds.	1607	100

823 AA 823 AA 823 AA 824 gil 824 gil 824 gil 825 gil 826 gil 826 gil 827 gil 827 gil 828 gil 829 g 829 g 830 g	No. 6453452 AY13402 AB18988 AB80270 13938181 6453540 10440436 7022318 7110152 17022600 19887215 17022560 17022033 18247250 AAB70772	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	mRNA; cDNA DKFZp434L0850 (from clone DKFZp434L0850). Amino acid sequence of protein PRO310. Amino acid sequence of a human transmembrane protein. Human PRO310 protein. , clone IMAGE:2905978, mRNA, partial cds. mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NTZRM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NTZRP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NTZRP2000239.	1607 1079 1079 2722 2455 807 1475 1219 592 315	Identity 100 63 63 63 99 99 100 100 80 98 43
823 AA 823 AA 823 AA 824 gil 824 gil 824 gil 825 gil 826 gil 826 gil 827 gil 827 gil 828 gil 828 gil 829 gil 829 gil 830 gil	AY13402 AB18988 AB80270 13938181 6453540 10440436 7022318 77110152 17022600 19887215 17022560 17022033 18247250	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	clone DKFZp434L0850). Amino acid sequence of protein PRO310. Amino acid sequence of a human transmembrane protein. Human PRO310 protein. , clone IMAGE:2905978, mRNA, partial cds. mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	1079 1079 2722 2455 807 1475 1219 592	63 63 63 99 99 100 100 80 98
823 AA 823 AA 824 gi 824 gi 824 gi 825 gi 826 gi 826 gi 827 gi 827 gi 828 A 828 gi 829 g 830 g 830 A 830 g	AB18988 AB80270 13938181 6453540 10440436 7022318 17110152 17022600 169887215 17022560 17022033 18247250	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	Amino acid sequence of a human transmembrane protein. Human PRO310 protein. , clone IMAGE:2905978, mRNA, partial cds. mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	1079 2722 2455 807 1475 1219 592	63 63 99 99 100 100 80 98
823 AA 823 AA 824 gi 824 gi 824 gi 825 gi 826 gi 826 gi 827 gi 827 gi 828 A 828 A 828 A 829 B 830 B 830 A 830 B	AB18988 AB80270 13938181 6453540 10440436 7022318 17110152 17022600 169887215 17022560 17022033 18247250	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	transmembrane protein. Human PRO310 protein. , clone IMAGE:2905978, mRNA, partial cds. mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	1079 2722 2455 807 1475 1219 592	63 99 99 100 100 80 98
824 gil 824 gic 824 gic 824 gi 825 gi 826 gi 826 gi 827 gi 827 gi 828 gi 828 A 828 g 829 g 829 g 830 g 830 A 830 g	13938181 6453540 10440436 7022318 77110152 17022600 19887215 17022560 17022033 18247250	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	Human PRO310 protein. , clone IMAGE:2905978, mRNA, partial cds. mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428): partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	2722 2455 807 1475 1219 592	99 99 100 100 80 98
824 git 824 git 824 git 824 git 824 git 825 git 826 git 826 git 827 git 827 git 828 git 828 A 828 git 829 g 830 g 830 A 830 g	13938181 6453540 10440436 7022318 77110152 17022600 19887215 17022560 17022033 18247250	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	, clone IMAGE:2905978, mRNA, partial cds. mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	2455 807 1475 1219 592 315	99 100 100 80 98
824 gi 825 gi' 826 gi 826 gi 827 gi 827 gi 827 gi 828 gi 828 A 828 gi 829 g 829 g 830 g	10440436 7022318 77110152 77022600 i9887215 i7022560 i7022033 i8247250	Homo sapiens Homo sapiens Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	mRNA; cDNA DKFZp434D0428 (from clone DKFZp434D0428); partial cds. mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	807 1475 1219 592	100 100 80 98
825 gi 826 gi 826 gi 826 gi 827 gi 827 gi 827 gi 828 gi 828 A 828 g 829 g 829 g 830 g 830 A 830 g	7022318 7110152 17022600 19887215 17022560 17022033 18247250	Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	mRNA for FLJ00053 protein, partial cds. cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	1475 1219 592 315	100 80 98
825 gi 826 gi 826 gi 826 gi 827 gi 827 gi 827 gi 828 gi 828 A 828 g 829 g 829 g 830 g 830 A 830 g	7022318 7110152 17022600 19887215 17022560 17022033 18247250	Mus musculus Homo sapiens Methanosarcin a thermophila Homo sapiens	cDNA FLJ10346 fis, clone NT2RM2001004. selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	1475 1219 592 315	100 80 98
826 gi 826 gi 826 gi 827 gi 827 gi 828 gi 828 A 828 g 829 g 829 g 830 g 830 A 830 g	i7022600 i9887215 i7022560 i7022033 i8247250	Methanosarcin a thermophila Homo sapiens	selenocysteine lyase SCLY cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	315	98
826 gi 826 gi 826 gi 827 gi 827 gi 828 gi 828 A 828 g 829 g 829 g 830 g 830 A 830 g	i7022600 i9887215 i7022560 i7022033 i8247250	Methanosarcin a thermophila Homo sapiens	cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	315	98
826 gi 827 gi 827 gi 828 gi 828 A 828 g 829 g 829 g 830 g 830 A 830 g	i9887215 i7022560 i7022033 i8247250	Methanosarcin a thermophila Homo sapiens	NT2RP2000764, weakly similar to NIFS PROTEIN. cysteine desulfurase NifS cDNA FLJ10491 fis, clone NT2RP2000239.	315	
827 gi 827 gi 828 gi 828 A 828 A 828 g 829 g 829 g 829 g 830 g	i7022560 i7022033 i8247250	a thermophila Homo sapiens	cDNA FLJ10491 fis, clone NT2RP2000239.		43
827 gi 828 gi 828 A 828 £ 829 £ 829 g 829 g 830 g	i7022033 i8247250	Homo sapiens	NT2RP2000239.	1266	
828 gi 828 A 828 g 829 g 829 g 829 g 830 g 830 A 830 g	i8247250	Homo sapiens			100
828 A 828 <u>g</u> 829 <u>g</u> 829 <u>g</u> 829 <u>g</u> 830 <u>A</u> 830 <u>g</u>			cDNA FLJ10156 fis, clone HEMBA1003447.	1161	97
828 g 829 g 829 g 829 g 830 g 830 A 830 g	AB70772	Homo sapiens	mRNA for neutral sphingomyelinase II (nSMase2 gene).	3489	100
829 g 829 g 829 g 830 g 830 A 830 g		Homo sapiens			100
829 g 829 g 829 g 830 g 830 A 830 g	i8247281	Mus musculus	neutral sphingomyelinase II	3187	91
829 g 829 g 830 g 830 A 830 g	17020945	Homo sapiens	cDNA FLJ20689 fis, clone KAIA2890.	2459	100
830 g 830 A 830 g	gi11596985	Homo sapiens	chromosome 14 clone RP11-361H10 map 14q24.3, complete sequence.	1819	97
830 A 830 g	gi6067151	Homo sapiens	chromosome 14 BAC 98L12, complete sequence.	1153	99
830 g	zi10039443	Homo sapiens	NEDL1 mRNA for NEDD4-like ubiquitin ligase 1, complete cds.	4335	56
830 g	A A 31/02 167	Homo sapiens	Human ZGGBP1 protein.	992	47
	AAW93167	Mus musculus	possible ubiquitin protein ligase	1062	50
	gi 1374782 gi 7021974	Homo sapiens	cDNA FLJ10115 fis, clone HEMBA1002777.	1882	99
		ļ	cDNA FLJ20739 fis, clone HEP07341.	1252	98
	gi7021027 gi5002381	Homo sapiens Takifugu	BAW	776	72
832 · §	gi7022523	rubripes Homo sapiens	cDNA FLJ10469 fis, clone NT2RP2000008, weakly similar to ZINC FINGER PROTEIN 84.	3772	99
832 §	gi1020145	Homo sapiens	Human DNA binding protein (HPF2) mRNA, complete cds.	1714	48
832 8	gi7243633	Homo sapiens	RB-associated KRAB repressor (RBAK) mRNA, complete cds.	1653	46
922	gi6433864	Homo sapiens	CLDN12 gene for claudin-12.	1295	100
	gi12053057	Homo sapiens	mRNA; cDNA DKFZp434I1816 (from clone DKFZp434I1816); complete cds.	1295	100
-	:0700000	Mus musculus		1125	91
	gi9799020 gi12053151	Homo sapiens	mRNA; cDNA DKFZp434G0326 (from	5605	99
			clone DKFZp434G0326); complete cds.	1268	88
	gi7020102 gi7023725	Homo sapiens Homo sapiens		719	100
835	R11073173	Homo sapiens	PLACE1007843. cDNA FLJ20583 fis, clone KAT09685.	2153	99

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
335	AAG02503	Homo sapiens	Human secreted protein, SEQ ID NO: 6584.	423	98
335	gi14289183	Homo sapiens	chac mRNA for chorein, complete cds.	193	24
336	gi7022600	Homo sapiens	cDNA FLJ10515 fis, clone NT2RP2000764, weakly similar to NIFS PROTEIN.		100
836	gi7110152	Mus musculus	selenocysteine lyase SCLY	1107	83
836	gi13592392	Caenorhabditis elegans	Contains similarity to Pfam domain: PF00266 (aminotran_5), Score=51.6, E- value=5.7e-12, N=1		44
837	gi7274380	Homo sapiens	group III secreted phospholipase A2 mRNA, complete cds.	2813	99
837	gi4314431	Homo sapiens	PAC clone RP3-412A9 from 22, complete sequence.	596	99
837	gi5627	Apis mellifera	phospholipase A-2	243	41
838	gi8331760	Homo sapiens	X28 region near ALD locus containing dual specificity phosphatase 9 (DUSP9), ribosomal protein L18a (RPL18a), Ca2+/Calmodulin-dependent protein kinase I (CAMKI), creatine transporter (CRTR), CDM protein (CDM), adrenoleukodystrophy protein (ALD), plexin-related protein (PLXB3), musclespecific serine kinase (MSSK), NADisocitrate dehydrogenase (IDH), translocon-associated protein delta (TRAP), and LU1 protein (LU1) genes, complete cds; and CCp pseudogene,		100
838	gi6651019	Mus musculus	complete sequence. semaphorin cytoplasmic domain-		50
838	gi6651021	Mus musculus			50
839	gi7023290	Homo sapiens	associated protein 3B cDNA FLJ10932 fis, clone OVARC1000588. 718		100
840	gi6094681	Homo sapiens	PAC clone RP5-1049N15 from 7q31.2-7q32, complete sequence.	4804	100
840	gi7264724	Homo sapiens	alpha-aminoadipate semialdehyde synthase mRNA, complete cds.	4804	100
840	gi4938304	Homo sapiens	mRNA for lysine-ketoglutarate reductase/saccharopine dehydrogenase, partial CDS.	4799	99
841	AAY66700	Homo sapiens	Membrane-bound protein PRO1137.	1164	95
841	AAB65223	Homo sapiens	Human PRO1137 (UNQ575) protein sequence SEQ ID NO:250.	1164	95
841	AAY50917	Homo sapiens	Human fetal brain cDNA clone vc4_1 derived protein.	1023	100
842	AAW56477	Homo sapiens	Amino acid sequence of human bone morphogenetic protein-16 (BMP-16).	1183	100
842	AAY03849	Homo sapiens	Human nodal protein.	1183	100
842	gi296605	Mus musculus		986	84
843	gi7020399	Homo sapiens		5470	100
843	gi10435659	Homo sapiens	cDNA FLJ13605 fis, clone PLACE1010562.	224	44
844	gi4886471	Homo sapiens	clone DKFZp586N0819).	531	100
845	gi3288470	Homo sapiens		728	100
845	gi3288452	Homo sapiens		334	94
845	gi3288468	Homo sapiens		334	94
846	gi14149050		turtle protein, isoform 4	1037	32

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		melanogaster			ļ
846	gi14149048	Drosophila melanogaster	turtle protein, isoform 3	1037	32
846	gi14149046	Drosophila melanogaster	turtle protein, isoform 2	939	34
847	gi7021049	Homo sapiens	cDNA FLJ20753 fis, clone HEP02714.	2930	99
847	gi9886896	Human herpesvirus 8	Orf73	175	20
847	gi11037008	Human herpesvirus 8	latent nuclear antigen	172	20
848	gi12597293	Homo sapiens	acidic mammalian chitinase precursor, mRNA, complete cds.	2018	100
848	gi6467177	Homo sapiens	TSA1902-L mRNA for novel member of chitinase family, complete cds.	2010	99
848	gi6467179	Homo sapiens	TSA1902-S mRNA for novel member of chitinase family, complete cds.	1725	99
849	gi32391	Homo sapiens	Human HOX4C mRNA for a homeobox protein.	1802	98
849	gi51416	Mus musculus	Hox-4.4	1591	88
849	gi4322104	Danio rerio	homeobox protein	425	82
850	gi1359443	Homo sapiens	Human gene for hepatitis C-associated microtubular aggregate protein p44, exon 9 and complete cds.	2299	99
850	AAY05371	Homo sapiens	Human HCMV inducible gene protein, SEQ ID NO 10.	2299	99
850	gi218576	Pan troglodytes	p44	2242	97
851	gi575494	Homo sapiens	MHC class II lymphocyte antigen beta- chain (HLA-DPB1a) mRNA, complete cds.	437	72
851	gi188479	Homo sapiens	Human MHC class II lymphocyte antigen (HLA-DP) beta chain mRNA, complete cds.	437	72
851	gi14044082	Homo sapiens	, Similar to major histocompatibility complex, class II, DP beta 1, clone MGC:14112, mRNA, complete cds.	429	70
852	gi181547	Homo sapiens	defensin 6 mRNA, complete cds.	318	90
852	AAR44819	Homo sapiens	Sequence of the gastrointestinal defensin (GID) peptide calledhuman defensin 6.	318	90
852	gi1200182	Homo sapiens	Human defensin 6 (HD-6) gene, complete cds.	314	89
853	gi13396914	Homo sapiens	The gene of C2GnT3	2389	100
853	gi7527464	Homo sapiens	core 2 beta-1,6-N- acetylglucosaminyltransferase 3 (C2GnT3) mRNA, complete cds.	2389	100
853	AAU00037	Homo sapiens	Human C2GnT3.	2389	100
855	gi7959772	Homo sapiens	PRO1483	252	100
856	gi5911169	Homo sapiens	transmembrane mucin 12 (MUC12) mRNA, partial cds.	2914	99
856	AAY59290	Homo sapiens	Human MUC12 polypeptide.	2914	99
856	gi2589172	Rattus norvegicus	mucin Muc3	595	36
857	AAE00508	Homo sapiens	Human lipase protein, MLip-1.	1456	100
857	gi56600	Rattus norvegicus	triacylglycerol lipase	776	58
857	gi3108175	Mus musculus	pancreatic lipase related protein 1	772	57
858	AAY94954	Homo sapiens	protein sequence SEQ ID NO:114.	1112	100
858	g:10434269	Homo sapiens	cDNA FLJ12650 fis, clone	872	100

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
	1,07		NT2RM4002054.		
858	gi7259234	Mus musculus	contains transmembrane (TM) region	660	60
859	gi7021851	Homo sapiens	cDNA FLJ10035 fis, clone HEMBA1000919.	1589	100
859	gi10440420	Homo sapiens	mRNA for FLJ00045 protein, partial cds.	654	89
859	AAY99671	Homo sapiens	Human GTPase associated protein-22.	654	89
860	gi7022523	Homo sapiens	Tidinan 617 ase assertant protest		99
860	gi1020145	Homo sapiens	Human DNA binding protein (HPF2) mRNA, complete cds.	1604	48
860	gi12584159	Homo sapiens	zinc finger protein 268 (ZNF268) mRNA, complete cds.	1542	48
861	gi6539434	Homo sapiens	SPR1 mRNA, complete cds.	808	100
861	gi6523547	Volvox carteri f. nagariensis	hydroxyproline-rich glycoprotein DZ- HRGP	185	39
861	gi904359	Beta vulgaris	chitinase 1	185	41
862	gi7021924	Homo sapiens	cDNA FLJ10081 fis, clone HEMBA1002018.	2742	100
862	gi10435862	Homo sapiens	cDNA FLJ13751 fis, clone PLACE3000339, weakly similar to GLUCOAMYLASE S1/S2 PRECURSOR (EC 3.2.1.3).	2687	99
862	gi11275988	Homo sapiens			99
863	gi7019913	Homo sapiens	cDNA FLJ20060 fis, clone COI.01358.	1830	100
863	gi10434817	Homo sapiens	cDNA FLJ13006 fis, clone NT2RP3000449.	1823	.99
863	gi10434659	Homo sapiens	cDNA FLJ12902 fis, clone NT2RP2004347.	1724	99
864	gi7329718	Homo sapiens			99
864	gi7022765	Homo sapiens	cDNA FLJ10619 fis, clone NT2RP2005472.	3153	99
864	gi14388939	Homo sapiens	chorea-acanthocytosis (CHAC) mRNA, complete cds.	462	30
865	gi28971	Homo sapiens	H.sapiens mRNA for autoantigen NOR- 90.	3813	100
865	gi509241	Homo sapiens	Human mRNA for upstream binding factor (hUBF).	2661	78
865	AAB44430	Homo sapiens	Human lung tumour-specific antigen encoded by cDNA	2649	78
866	gi13445482	Homo sapiens	HP43.8KD mRNA, complete cds.	282	47
866	gi10434108	Homo sapiens	cDNA FLJ12552 fis, clone NT2RM4000712, moderately similar to Homo sapiens ubiquitin hydrolyzing enzyme I (UBH1) mRNA.	219	36
866	gi10436670	Homo sapiens	cDNA FLJ14256 fis, clone PLACE1000007, weakly similar to PROBABLE UBIQUITIN CARBOXYL- TERMINAL HYDROLASE R10E11.3 (EC 3.1.2.15).	219	36
967	A A D 77770	Uomo caniere	Human phosphatase MTMR7_h.	743	57
867	AAB73229 gi5901814	Homo sapiens Drosophila melanogaster	BcDNA.GH04637	503	48
867	gi7020021	Homo sapiens	cDNA FLJ20126 fis, clone COL06160.	697	73
868	g:7959801	Homo sapiens	PRO0800	392	100
869	gi12654971	Homo sapiens	, calcium-regulated heat-stable protein (24kD), clone MGC:5586, mRNA,	417	97

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			complete cds.		
869	gi13097198	Homo sapiens	, calcium-regulated heat-stable protein (24kD), clone MGC:5235, mRNA, complete cds.	417	97
869	AAW61023	Homo sapiens	Human RNA binding protein.	417	97
870	gi6650832	Homo sapiens	PRO2086	243	100
871	gi2217942	Rattus norvegicus	glycoprotein specific UDP- glucuronyltransferase	1802	97
871	gi8051678	Homo sapiens	hu-GlcAT-P mRNA for glucuronyltransferase, complete cds.	1757	99
871	gi4519214	Rattus norvegicus	UDP-glucuronyltransferase-S	760	50
872	gi14286288	Homo sapiens	, Similar to RIKEN cDNA 2010004P11 gene, clone MGC:2734, mRNA, complete cds.	715	100
872	gi13529665	Mus musculus	RIKEN cDNA 2010004P11 gene	706	98
872	gi2565364	Musca domestica	Sex-lethal protein	134	33
873	gi190406	Homo sapiens	Human profilaggrin gene exons 1-3, 5' end.	6301	99
873	gi190396	Homo sapiens	Human profilaggrin gene, 3' end.	5133	99
873	gi190404	Homo sapiens	Human profilaggrin mRNA, 3' end.	3696	89
874	gi791002	Homo sapiens	ARSD gene, complete CDS.	1761	99
874	gi6651286	Homo sapiens	arylsulfatase D beta (ARSD) mRNA, complete cds.		99
874	gi791004	Homo sapiens	ARSE gene, complete CDS.	947	58
875	gi13097675	Homo sapiens	protein HCDASE, clone MGC:1171, mRNA, complete cds.		96
875	AAY87599	Homo sapiens	Human fatty acid beta-oxidation enzyme 6 HUFA-2.		96
875	AAG03352	Homo sapiens	Human secreted protein, SEQ ID NO: 7433.	591	100
876	gi6180180	Homo sapiens	transcription factor IGHM enhancer 3, JM11 protein, JM4 protein, JM5 protein, T54 protein, JM10 protein, A4 differentiation-dependent protein, triple LIM domain protein 6, and synaptophysin genes, complete cds; and L-type calcium channel alpha-1 subunit gene, partial cds, complete sequence.	908	100
876	gi3114826	Homo sapiens	mRNA for IM4 protein, complete CDS (clone IMAGE 546750 and LLNLc110F1857Q7 (RZPD Berlin)).	908	100
876	gi7673612	Mus musculus	DXImx39e protein	831	91
877	gi13543663	Homo sapiens	, ubiquitin-conjugating enzyme E2D l (homologous to yeast UBC4/5), clone MGC:14673, mRNA, complete cds.	805	100
877	gi460810	Homo sapiens	H.sapiens UBCH5 mRNA for ubiquitin conjugating enzyme.	805	100
877	gi4868140	Homo sapiens	ubiquitin-conjugating enzyme HBUCE1 mRNA, complete cds.	747	91
878	gi7020915	Homo sapiens	cDNA FLJ20666 fis, clone KAIA608.	1288	100
878	gi3169096	Schizosacchar omyces pombe	possible pre-mRNA processing by similarity to yeast prp39	279	33
878	gi10177721	Arabidopsis thaliana	gene_id:MPL12.20~	146	22
879	gi7020681	Homo sapiens	cDNA FLJ20519 fis, clone KAT10365.	891	100
879	AAY87267	Home sapiens	Human signal peptide containing protein	824	95

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			HSPP-44 SEQ ID NO:44.		ļ <u> </u>
879	AAB65245	Homo sapiens	Human PRO1104 (UNQ547) protein sequence SEQ ID NO:297.	824	95
880	gi6560622	Homo sapiens	PRO0611	501	100
881	AAB57079	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1657.	668	100
881	AAY99372	Homo sapiens	Human PRO1430 (UNQ736) amino acid sequence SEQ ID NO:116.	668	100
881	AAB88356	Homo sapiens	Human membrane or secretory protein clone PSEC0082.	661	99
882	gi1381181	Oryctolagus cuniculus	ubiquitin-conjugating enzyme E2-32k	663	100
882	gi13436071	Homo sapiens	, clone MGC:10481, mRNA, complete cds.	663	100
882	gi7020506	Homo sapiens	cDNA FLJ20419 fis, clone KAT02435.	658	99
883	gi1381181	Oryctolagus cuniculus	ubiquitin-conjugating enzyme E2-32k	1265	99
883	gi13436071	Homo sapiens	, clone MGC:10481, mRNA, complete cds.	1265	99
883	gi7020506	Homo sapiens	cDNA FLJ20419 fis, clone KAT02435.	1256	98
884	gi1381181	Oryctolagus cuniculus	ubiquitin-conjugating enzyme E2-32k	383	97
884	gi13436071	Homo sapiens	, clone MGC:10481, mRNA, complete cds.	383	97
884	gi7020506	Homo sapiens	cDNA FLJ20419 fis, clone KAT02435.	383	97
885	gi14424536	Homo sapiens	, Similar to septin 6, clone MGC:16619, mRNA, complete cds.	2183	99
885	gi5689158	Mus musculus	Septin6	2114	95
885	gi7023141	Homo sapiens	cDNA FLJ10849 fis, clone NT2RP4001414, highly similar to SEPTIN 2 HOMOLOG.	1840	82
886	gi14424536	Homo sapiens	, Similar to septin 6, clone MGC:16619, mRNA, complete cds.	1213	63
886	gi5689158	Mus musculus	Septin6	1162	62
886	gi7023141	Homo sapiens	cDNA FLJ10849 fis, clone NT2RP4001414, highly similar to SEPTIN 2 HOMOLOG.	995	51
887	gi4309951	Homo sapiens	BAC clone RP11-121A8 from 7p14-p13, complete sequence.	684	100
887	AAG00417	Homo sapiens	Human secreted protein, SEQ ID NO: 4498.	684	100
887	gi339159	Homo sapiens	Human T-cell receptor germline gamma- chain gene V-region (V3; subgroup I).	392	73
888	gi2570015	Homo sapiens	H.sapiens PAX7 gene, exon 1 (and joined CDS).	2756	100
888	gi2570021	Homo sapiens	H.sapiens mRNA for paired box containing transcription factor, PAX7.	2756	100
888	gi2570014	Homo sapiens	H.sapiens PAX7 gene, exon 1 (and joined CDS).	2735	99

TABLE 3

IABLE 3		Description	Results*
SEQ ID NO:	Accession No.	Description	
445	BL00434	HSF-type DNA-binding domain proteins.	BL00434C 23.85 7.111e-09 1089-1129
446	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 1.000e-13 216-229 PD00066 13.92 2.286e-12 244-257 PD00066 13.92 4.522e-11 299-312
			PD00066 13.92 6.538e-10 157-170 PD00066 13.92 7.923e-10 327-340
453	PR00037	LACR BACTERIAL REGULATORY PROTEIN HTH SIGNATURE	PR00037A 12.66 6.786e-09 34-49
465	PR00320	G-PROTEIN BETA WD-40 REPEAT SIGNATURE	PR00320C 13.01 6.100e-09 197-212 PR00320C 13.01 6.400e-09 393-408 PR00320A 16.74 8.683e-09 197-212 PR00320B 12.19 9.775e-09 299-314
466	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 5.881e-09 14-47
470	BL00175	Phosphoglycerate mutase family phosphohistidine proteins.	BL00175D 27.67 8.500e-40 175-227 BL00175C 23.75 5.000e-25 90-122 BL00175A 15.42 8.333e-20 17-37 BL00175B 12.60 1.000e-12 66-79
472	BL00315	Dehydrins proteins.	BL00315A 9.35 8.119e-09 105-133
473	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 4.000e-11 44-53
475	PD02448	TRANSCRIPTION PROTEIN DNA-BINDIN.	PD02448A 9.37 4.293e-09 171-210
477	PR00625	DNAJ PROTEIN FAMILY SIGNATURE	PR00625A 12.84 8.500e-19 121-141 PR00625B 13.48 3.204e-15 151-172
478	PD02102	SUBUNIT E V-ATPASE VACUOLAR ATP SYNTHASE HYDROL.	PD02102A 16.74 5.853e-10 26-70
479	BL00018	EF-hand calcium-binding domain proteins.	BL00018 7.41 4.706e-11 49-62
480	PR00501	KELCH REPEAT SIGNATURE	PR00501A 8.25 9.182e-09 544-558
483	PR00878	CHOLINESTERASE SIGNATURE	PR00878F 5.37 5.179e-12 500-513
484	BL00378	Hexokinases proteins.	BL00378C 16.14 1.000e-40 207-251 BL00378E 22.92 1.000e-40 725-771
			BL00378C 16.14 3.520e-40 655-699 BL00378E 22.92 3.382e-36 277-323 BL00378B 14.23 5.333e-35 509-546 BL00378B 14.23 8.953e-28 61-98
			BL00378A 19.01 1.346e-22 22-50 BL00378F 8.27 2.688e-17 893-908 BL00378D 10.94 6.294e-17 703-715 BL00378D 10.94 5.500e-16 255-267
			BL00378F 8.27 9.609e-13 445-460 BL00378A 19.01 3.017e-12 470-498
485	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 2.688e-15 352-369 BL00028 16.07 4.375e-15 324-341 BL00028 16.07 4.176e-14 604-621
			BL00028 16.07 8.412e-14 380-397 BL00028 16.07 9.471e-14 576-593
			BL00028 16.07 1.450e-13 548-565 BL00028 16.07 2.350e-13 436-453
			BL00028 16.07 4.150e-13 492-509 BL00028 16.07 5.050e-13 296-313
1			BL00028 16.07 1.783e-12 520-537

SEQ ID NO:	Accession No.	Description	Results*
NO:	110.		BL00028 16.07 3.348e-12 632-649
			BL00028 16.07 5.304e-12 408-425
			BL00028 16.07 5.304e-12 660-677
			BL00028 16.07 4.808e-11 464-481
	ĺ		BL00028 16.07 7.000e-10 268-285
486	BL00301	GTP-binding elongation	BL00301B 20.09 1.429e-26 128-160
400	BLOOJOI	factors proteins.	BL00301A 12.41 6.400e-15 62-74
487	PD00301	PROTEIN REPEAT	PD00301B 5.49 7.600e-12 826-837
407	1.000301	MUSCLE CALCIUM-BI.	
489	BL00227	Tubulin subunits alpha, beta,	BL00227B 19.29 1.000e-40 52-107
409	BL00227	and gamma proteins.	BL00227C 25.48 1.000e-40 113-165
		and gamma proteins.	BL00227D 18.46 1.000e-40 222-276
			BL00227F 21.16 1.000e-40 382-436
			BL00227E 24.15 6.727e-36 326-361
			BL00227A 24.55 2.125e-33 1-35
100	DI 00470	Phorbol esters /	BL00479B 12.57 6.625e-09 1271-1287
490	BL00479		BE00479B 12.57 0.025C-05 1271-1207
		diacylglycerol binding	
	DI 00 170	domain proteins.	BL00479B 12.57 6.625e-09 1250-1266
491	BL00479	Phorbol esters /	DLUG4/7D 12.37 0.0236-07 1230-1200
		diacylglycerol binding	
		domain proteins.	BL00107A 18.39 5,500e-19 138-169
492	BL00107	Protein kinases ATP-binding	¹ BL00107B 13.31 1.000e-16 203-219
		region proteins.	BL50002A 14.19 5.000e-15 392-411
493	BL50002	Src homology 3 (SH3)	
		domain proteins profile.	BL50002B 15.18 2.500e-09 430-444
494	PR00049	WILM'S TUMOUR	PR00049D 0.00 6.949e-09 87-102
		PROTEIN SIGNATURE	
497	BL00914	Syntaxin / epimorphin	BL00914 24.91 6.172e-09 249-299
		family proteins.	· · · · · · · · · · · · · · · · · · ·
498	PD00066	PROTEIN ZINC-FINGER	PD00066 13.92 8.200e-16 362-375
		METAL-BINDI.	PD00066 13.92 4.462e-15 334-347
			PD00066 13.92 8.615e-15 473-486
			PD00066 13.92 5.200e-14 306-319
			PD00066 13.92 3.000e-13 390-403
500	PF00780	Domain found in NIK1-like	PF007801 14.69 7.863e-09 293-323
		kinases, mouse citron and	
		yeast ROM.	
501	BL00518	Zinc finger, C3HC4 type	BL00518 12.23 7.333e-09 279-288
<u></u>		(RING finger), proteins.	-
502	DM01418	352 FIBRILLAR	DM01418A 20.83 2.050e-23 1537-1585
	Ī	COLLAGEN CARBOXYL-	DM01418B 22.51 5.895e-21 1632-1674
		TERMINAL.	DM01418C 20.48 8.571e-18 1702-1744
508	BL01052	Calponin family repeat	BL01052B 15.31 1.000e-09 131-157
ļ		proteins.	
512	BL01310	ATPIG1/PLM/MAT8	BL01310 14.74 7.107e-36 27-63
		family proteins.	
515	DM00475	w LOW TRANSPOSASE	DM00475B 12.12 6.019e-09 386-406
ļ		SAPA 12K.	ļ
516	BL00636	Nt-dnaJ domain proteins.	BL00636A 8.07 5.865e-11 64-81
519	PR00625	DNAJ PROTEIN FAMILY	PR00625A 12.84 2.019e-14 76-96
	1	SIGNATURE	PR00625B 13.48 5.714e-11 106-127
520	BL00216	Sugar transport proteins.	BL00216B 27.64 6.400e-10 92-142
523	BL01033	Globins profile.	BL01033B 13.81 1.000e-15 38-50
526	BL50002	Src homology 3 (SH3)	BL50002B 15.18 4.750e-12 1075-1089
320	DD50002	domain proteins profile.	
531	PR00249	SECRETIN-LIKE GPCR	PR00249G 15.72 8.892e-10 387-409
221	F KUU249	SUPERFAMILY	PR00249C 17.08 6.609e-09 223-247
		SIGNATURE	11002470 17.00 0.0070 07 223 247
L	BL00528	Ribosomal protein S4e	BL00528D 27.17 8.012e-09 341-395
532			

SEQ ID NO:	Accession No.	Description	Results*
534	PR00194	TROPOMYOSIN SIGNATURE	PR00194C 6 38 1.900e-35 109-138 PR00194E 8.74 1.000e-30 195-221
			PR00194D 9.57 8.714e-27 139-163 PR00194B 10.24 2.800e-25 84-105 PR00194A 7.86 5.500e-22 48-66
535	PR00194	TROPOMYOSIN SIGNATURE	PR00194C 6.38 1.900e-35 109-138 PR00194E 8.74 1.000e-30 195-221 PR00194B 10.24 2.800e-25 84-105
			PR00194D 9.57 1.900e-23 139-163 PR00194A 7.86 5.500e-22 48-66
538	PR00019	LEUCINE-RICH REPEAT SIGNATURE	PR00019A 11.19 5.050e-11 110-124
541	BL00540	Ferritin iron-binding regions proteins.	BL00540A 15.06 1.000e-40 32-73 BL00540B 18.82 1.000e-40 123-178 BL00540C 13.00 7.750e-14 188-200
546	PR00153	CYCLOPHILIN PEPTIDYL-PROLYL CIS- TRANS ISOMERASE SIGNATURE	PR00153E 9.10 2.385e-15 121-137
548	BL00115	Eukaryotic RNA polymerase II heptapeptide repeat proteins.	BL00115Z 3.12 8.213e-09 63-112
549	BL01282	BIR repeat proteins.	BL01282B 30.49 2.373e-12 317-356
551	BL00570	Bacterial ring hydroxylating dioxygenases alpha-subunit signa.	BL00570B 19.03 9.357e-09 277-309
553	PD01427	TRANSFERASE METHYLTRANSFERASE BI.	PD01427B 22.45 7.000e-11 127-168
554	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048A 10.52 7.632e-11 447-461
555	PD02637	SERUM PARAOXONASE/ARYLES TERASE P.	PD02637A 14.26 1.000e-40 32-87 PD02637G 13.82 1.000e-40 307-355 PD02637D 13.69 6.053e-36 170-218 PD02637B 10.33 8.875e-34 106-141 PD02637E 11.92 8.200e-28 218-249 PD02637C 7.53 3.520e-27 141-170 PD02637F 15.62 9.438e-26 281-307
556	DM00892	3 RETROVIRAL PROTEINASE.	DM00892C 23.55 2.768e-16 474-508
557	BL00039	DEAD-box subfamily ATP-dependent helicases proteins.	BL00039D 21.67 5.179c-36 294-340 BL00039A 18.44 7.955c-29 15-54 BL00039C 15.63 1.300c-16 143-167 BL00039B 19.19 2.465c-12 58-84
558	PR00507	N12 CLASS N6 ADENINE- SPECIFIC DNA METHYLTRANSFERASE SIGNATURE	PR00507B 14.16 8.932e-09 83-98
559	BL00383	Tyrosine specific protein phosphatases proteins.	BL00383E 10.35 8.683e-12 242-253
566	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 5.500e-13 214-227
572	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 4.432e-09 76-130
573	BL00422	Granins proteins.	BL00422C 16.18 4.638e-10 49-77
574	PR00319	BETA G-PROTEIN (TRANSDUCIN) SIGNATURE	PR00319A 15.27 7.911e-10 452-469 PR00319A 15.27 2.180e-09 410-427
577	BL00269	Mammalian defensins proteins.	BL00269C 16.52 6.786e-26 73-102 BL00269A 8.53 2.607e-20 8-28 BL00269B

SEQ ID NO:	Accession No.	Description	Results*
			19.17 5.500e-17 35-64
578	PD02327	GLYCOPROTEIN ANTIGEN PRECURSOR IMMUNOGLO.	PD02327B 19.84 2.241e-11 157-179
579	BL00596	High potential iron-sulfur proteins.	BL00596B 13.07 9.743e-09 273-285
580	BL00915	Phosphatidylinositol 3- and 4-kinases proteins.	BL00915C 22.43 8.147e-32 1015-1054 BL00915D 27.02 9.217e-27 1092-1128 BL00915B 22.78 3.382e-25 918-956 BL00915A 10.09 5.500e-10 756-768
584	BL00038	Myc-type, 'helix-loop-helix' dimerization domain proteins.	BL00038B 16.97 7.488e-09 499-520
585	BL00795	Involucrin proteins.	BL00795C 17.06 9.200e-09 498-543
586	BL00710	Phosphoglucomutase and phosphomannomutase phosphoserine signa.	BL00710 12.98 9.100e-17 159-174
587	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.714e-10 34-43
588	PR00326	GTP1/OBG GTP-BINDING PROTEIN FAMILY SIGNATURE	PR00326A 8.75 5.979e-14 257-278
591	BL00548	Ribosomal protein S3 proteins.	BL00548 20.58 7.000e-19 66-96
592	BL00478	LIM domain proteins.	BL00478B 14.79 1.250e-12 557-572 BL00478B 14.79 6.000e-12 494-509 BL00478B 14.79 2.400e-11 624-639
594	PR00109	TYROSINE KINASE CATALYTIC DOMAIN SIGNATURE	PR00109B 12.27 3.681e-13 141-160
596	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 9.063e-12 510-525 PR00049D 0.00 8.286e-10 513-528 PR00049D 0.00 9.000e-10 509-524 PR00049D 0.00 9.429e-10 511-526
599	BL00232	Cadherins extracellular repeat proteins domain proteins.	BL00232B 32.79 4.750e-40 142-190 BL00232A 27.72 3.793e-22 48-81 BL00232B 32.79 1.257e-16 251-299 BL00232C 10.65 5.935e-14 249-267 BL00232D 16.25 3.368e-13 763-778 BL00232B 32.79 3.512e-11 366-414
600	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 9.695e-09 513-546
601	PF00583	Acetyltransferase (GNAT) family.	PF00583B 10.18 9.100e-10 120-130
602	PR00326	GTP1/OBG GTP-BINDING PROTEIN FAMILY SIGNATURE	PR00326A 8.75 5.950e-11 146-167
604	BL00319	Amyloidogenic glycoprotein extracellular domain proteins.	
607	BL00239	Receptor tyrosine kinase class II proteins.	BL00239F 28.15 4.717e-25 477-522 BL00239E 17.14 5.897e-23 423-473 BL00239C 18.75 7.600e-17 372-395
608	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 3.357e-32 10-49
609	PR00449	TRANSFORMING PROTEIN P21 RAS SIGNATURE	PR00449A 13.20 4.808e-10 5-27 PR00449D 10.79 5.636e-09 111-125
610	PF00791	Domain present in ZO-1 and	PF00791C 20.98 2.412e-09 1-40
		TRANSFORMING PROTEIN P21 RAS SIGNATURE	PR00449D 10.79 5.636e-09 111-125

SEQ ID NO:	Accession No.	Description	Results*
612	PR00109	Unc5-like netrin receptors. TYROSINE KINASE CATALYTIC DOMAIN SIGNATURE	PR00109B 12.27 9.234e-13 487-506
613	BL00678	Trp-Asp (WD) repeat proteins proteins.	BL00678 9.67 1.600e-10 104-115 BL00678 9.67 5.737e-09 62-73 BL00678 9.67 8.105e-09 146-157 BL00678 9.67 8.105e-09 276-287
615	PR00334	HMW KININOGEN SIGNATURE	PR00334B 8.69 5.230e-10 460-484 PR00334B 8.69 1.771e-09 464-488 PR00334B 8.69 2.886e-09 466-490 PR00334B 8.69 8.200e-09 458-482
617	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 5.881e-09 66-99
618	PF00084	Sushi domain proteins (SCR repeat proteins.	PF00084B 9.45 7.188e-10 539-551 PF00084B 9.45 7.300e-09 600-612
619	PR00169	POTASSIUM CHANNEL SIGNATURE	PR00169A 16.77 4.316e-09 72-92
621	BL00845	CAP-Gly domain proteins.	BL00845 16.43 1.900e-25 321-346 BL00845 16.43 9.325e-22 443-468
622	BL01002	Translationally controlled tumor protein.	BL01002D 18.24 4.706e-26 143-171 BL01002C 21.97 6.143e-26 79-110 BL01002A 13.19 1.360e-24 1-24 BL01002B 7.39 3.118e-14 48-62
624	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 8.857e-10 1030-1045
627	PR00011	TYPE III EGF-LIKE SIGNATURE	PR00011A 14.06 4.822e-09 475-494
629	PF00930	Dipeptidyl peptidase IV (DPP IV) N-terminal region.	PF009301 15.96 5.000e-15 656-684 PF00930J 8.78 6.045e-12 708-729
630	PF00930	Dipeptidyl peptidase IV (DPP IV) N-terminal region.	PF00930I 15.96 5.000e-15 598-626 PF00930J 8.78 6.045e-12 650-671
631	BL00303	S-100/ICaBP type calcium binding protein.	BL00303B 26.15 1.844e-10 365-402
632	BL00114	Phosphoribosyl pyrophosphate synthetase proteins.	BL00114A 17.22 1.000e-40 54-101 BL00114B 15.90 1.000e-40 107-153 BL00114D 21.45 1.000e-40 208-259 BL00114C 18.22 2.895e-34 167-202 BL00114E 14.48 3.647e-25 293-317
635	BL00870	Chaperonins clpA/B proteins.	BL00870F 8.73 4.833e-36 376-425 BL00870G 8.07 6.553e-27 436-470 BL00870E 17.62 3.333e-16 304-359
639	BL00633	Bromodomain proteins.	BL00633B 13.82 9.775e-13 237-262 BL00633B 13.82 4.750e-11 80-105
641	BL00299	Ubiquitin domain proteins.	BL00299 28.84 7.962e-17 47-99
642	PD02102	SUBUNIT E V-ATPASE VACUOLAR ATP SYNTHASE HYDROL.	PD02102A 16.74 4.176e-10 97-141
643	PD02080	T-CELL GLYCOPROTEIN CD8 CHAIN SURFACE ALPHA PRE.	PD02080D 15.22 6.557e-09 269-306
644	BL01245	RIO1/ZK632.3/MJ0444 family proteins.	BL01245F 18.75 7.805e-14 239-276
646	BL00469	Nucleoside diphosphate kinases proteins.	BL00469 22.22 1.000e-40 41-96
649	PR00217	43 KD POSTSYNAPTIC PROTEIN SIGNATURE	PR00217C 10.91 5.945e-09 91-107
651	PR00326	GTP1/OBG GTP-BINDING PROTEIN FAMILY SIGNATURE	PR00326A 8.75 7 600e-11 629-650

SEQ ID NO:	Accession No.	Description	Results*
652	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 8.322e-09 227-260
653	PF01298	Transferrin binding protein.	PF01298C 15.13 1.000e-08 413-440
658	PR00443	G-PROTEIN ALPHA SUBUNIT GROUP S SIGNATURE	PR00443A 15.16 9.451e-09 89-105
659	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.714e-10 34-43
663	BL00466	TFIIS zinc ribbon domain proteins.	BL00466 25.88 1.000e-32 294-331
664	PD00567	PROTEIN RNA-BINDING RNA REPEAT HYD.	PD00567B 18.23 3.172e-10 411-425
665	BL00030	Eukaryotic RNA-binding region RNP-1 proteins.	BL00030A 14.39 7.882e-11 10-29
669	PR00124	ATP SYNTHASE C SUBUNIT SIGNATURE	PR00124A 8.81 8.347e-11 117-137
670	PD01234	PROTEIN NUCLEAR BROMODOMAIN TRANS.	PD01234B 15.53 2.500e-10 38-56
671	BL00466	TFIIS zinc ribbon domain proteins.	BL00466 25.88 1.000e-32 219-256
672	BL01282	BIR repeat proteins.	BL01282B 30.49 2.068e-12 298-337
673	BL00455	Putative AMP-binding domain proteins.	BL00455 13.31 4.176e-14 201-217
674	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 8.703e-10 407-461 BL01160B 19.54 2.373e-09 414-468
675	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.286e-10 326-335
676	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 5.286e-10 335-344
682	PR00761	BINDIN PRECURSOR SIGNATURE	PR00761E 14.32 4.789e-09 499-518
691	BL00415	Synapsins proteins.	BL00415Q 2.23 2.885e-09 83-119
692	PR00211	GLUTELIN SIGNATURE	PR00211B 0.86 6.167e-09 115-136
694	PR00320	G-PROTEIN BETA WD-40 REPEAT SIGNATURE	PR00320C 13.01 7.300e-09 297-312
696	PD02952	KINASE TRANSFERASE CHOLINE PROTEIN MULTIGENE FAMI.	PD02952C 15.76 5.701e-16 263-293 PD02952B 15.57 7.242e-11 243-257 PD02952A 11.84 9.625e-09 131-159
697	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 7.231e-15 504-517 PD00066 13.92 5.800e-14 220-233 PD00066 13.92 1.000e-11 248-261 PD00066 13.92 5.696e-11 333-346 PD00066 13.92 2.500e-09 361-374
698	PR00205	CADHERIN SIGNATURE	PR00205B 11.39 6.571e-13 167-185
699	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 4.966e-09 50-65 PR00049D 0.00 9.237e-09 74-89
701	PR00988	URIDINE KINASE SIGNATURE	PR00988A 6.39 6.600e-15 98-116 PR00988C 13.64 5.605e-13 175-191 PR00988E 8.27 8.393e-13 245-257 PR00988D 5.95 8.250e-11 231-242 PR00988F 12.23 9.820e-11 267-281 PR00988B 11.60 2.317e-10 128-140
702	PR00625	DNAJ PROTEIN FAMILY SIGNATURE	PR00625A 12.84 1.804e-13 22-42 PR00625B 13.48 5.821e-13 53-74
706	PF00023	Ank repeat proteins.	PF00023A 16.03 2.286e-09 209-225
708	BL01212	ATP P2X receptors proteins.	BL01212A 34.89 1.000e-40 43-96 BL01212E 24.87 1.000e-40 227-282 BL01212D 11.42 6.700e-25 185-209 BL01212G 11.86 2.800e-24 310-338 BL01212B 19.25 3.571e-21 129-154

SEQ ID NO:	Accession No.	Description	Results*
NO:	110.		BL01212C 8.40 1.214c-14 162-173 BL01212F 10.12 4.774e-14 291-302
700	BL00194	Thioredoxin family proteins.	BL00194 12.16 3.455e-17 45-58
709	BL00194 BL00439	Acyltransferases ChoActase	BL00439F 26.22 1.000e-40 418-471
712	BL00439	/ COT / CPT family	BL00439E 19.05 2.440e-24 320-349
	1	proteins.	BL00439B 16.82 1.000e-20 167-189
	1	proteins.	BL00439H 18.24 4.600e-20 566-592
			BL00439A 9.40 1.237e-15 35-52
			BL00439D 13.11 4.545e-15 272-290
			BL00439C 13.53 1.730e-11 248-261
	1		BL00439G 13.40 9.719e-11 513-524
	DI 00412	Neuromodulin (GAP-43)	BL00412D 16.54 8.990e-09 305-356
716	BL00412	proteins.	
718	BL01271	Sodium:sulfate symporter	BL01271D 25.26 5.979e-32 537-592
:		family proteins.	BL01271A 8.06 6.250e-18 131-151
			BL01271C 13.62 7.750e-17 464-486
	1		BL01271B 12.02 1.563e-16 269-294
719	PF00023	Ank repeat proteins.	PF00023B 14.20 2.500e-10 141-151
			PF00023A 16.03 4.000e-10 112-128
721	PF00023	Ank repeat proteins.	PF00023A 16.03 1.750e-10 66-82
			PF00023B 14.20 5.500e-09 161-171
1	ļ		PF00023A 16.03 8.714e-09 363-379
725	PR00019	LEUCINE-RICH REPEAT	PR00019B 11.36 1.500e-11 173-187
}		SIGNATURE	PR00019A 11.19 2.800e-11 314-328
1		1	PR00019A 11.19 5.050e-11 176-190
†			PR00019B 11.36 3.520e-09 311-325
.			PR00019B 11.36 4.600e-09 541-555
			PR00019B 11.36 5.320e-09 471-485
	1		PR00019A 11.19 6.000e-09 544-558
1		<u> </u>	PR00019B 11.36 8.200e-09 242-256
_			PR00019B 11.36 9.640e-09 127-141
731	PR00681	RIBOSOMAL PROTEIN SI SIGNATURE	PR00681[8.81 9.897e-09 600-619
736	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL-	PD01066 19.43 9.581e-31 8-47
1		BINDING NU.	
739	BL00972	Ubiquitin carboxyl-terminal	BL00972A 11.93 1.587e-13 170-188
		hydrolases family 2 proteins.	BL00972D 22.55 8.826e-11 590-615
740	BL00972	Ubiquitin carboxyl-terminal	BL00972A 11.93 1.587e-13 170-188
		hydrolases family 2 proteins.	BL00972D 22.55 8.826e-11 590-615
741	DM01688	2 POLY-IG RECEPTOR.	DM01688G 16.45 6.936e-10 85-117
747	PF00646	F-box domain proteins.	PF00646A 14.37 6.625e-09 50-64
753	BL01168	Ribosomal protein S27e	BL01168 15.74 1.000e-40 20-75
756	PD00066	PROTEIN ZINC-FINGER	PD00066 13.92 6.885e-10 127-140
756	ססטטטערן	METAL-BINDI.	120000 13.72 0.0000 10 12.7
	DD00201	PROTEIN REPEAT	PD00301B 5.49 7.231e-09 1019-1030
757	PD00301	MUSCLE CALCIUM-BI.	
761	BL00712	Ribosomal protein S17e	BL00712B 12.56 1.000e-40 28-66
		proteins.	BL00712A 6.23 8.855e-19 2-22
762	PF00878	Cation-independent mannose-6-phosphate	PF00878T 17.51 3.818e-09 799-826
		receptor repeat proteins.	
763	BL00303	S-100/ICaBP type calcium	BL00303A 21.77 9.526e-31 3-40
103	DEGGGG	binding protein.	BL00303B 26.15 5.737e-30 50-87
766	BL00018	EF-hand calcium-binding	BL00018 7.41 6.087e-09 237-250
. 700	DEUUUIA	domain proteins.	
768	BL00221	MIP family proteins.	BL00221D 12.33 6.143e-19 180-195
700	5200221		BL00221C 13.36 1.000e-14 135-152
	1)	BL00221E 8.47 3.739e-13 203-214

SEQ ID NO:	Accession No.	Description	Results*
			BL00221B 10.22 1.750e-12 63-74 BL00221A 6.39 5.200e-12 16-27
769	PF00992	Troponin.	PF00992A 16.67 8.859e-10 214-249
770	BL00720	Guanine-nucleotide dissociation stimulators CDC25 family sign.	BL00720B 16.57 8.297e-15 136-160
771	PR00883	HIGH MOBILITY GROUP- LIKE NUCLEAR PROTEIN SIGNATURE	PR00883A 6.49 8.920e-09 191-205
772	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 6.786e-32 8-47
775	BL00301	GTP-binding elongation factors proteins.	BL00301B 20.09 5.500e-31 90-122 BL00301C 11.73 8.200e-15 423-437 BL00301A 12.41 3.842e-13 9-21
776	PR00453	VON WILLEBRAND FACTOR TYPE A DOMAIN SIGNATURE	PR00453A 12.79 4.892e-12 325-343 PR00453B 14.65 1.614e-10 162-177 PR00453A 12.79 3.152e-10 123-141
779	PR00399	SYNAPTOTAGMIN SIGNATURE	PR00399A 9.52 1.730e-13 145-161 PR00399B 14.27 2.059e-13 160-174 PR00399C 12.82 7.324e-12 216-232 PR00399D 14.48 3.930e-10 236-247 PR00399B 14.27 1.915e-09 291-305
780	BL00115	Eukaryotic RNA polymerase II heptapeptide repeat proteins.	BL00115Z 3.12 8.395e-10 123-172 BL00115Z 3.12 4.375e-09 137-186
783	PD00066	PROTEIN ZINC-FINGER METAL-BINDI.	PD00066 13.92 8.800e-14 165-178 PD00066 13.92 8.800e-14 193-206 PD00066 13.92 5.286e-12 249-262 PD00066 13.92 8.269e-10 221-234
786	PF00975	Thioesterase domain proteins.	PF00975B 10.82 2.688e-12 90-104
788	DM01970	0 kw ZK632.12 YDR313C ENDOSOMAL III.	DM01970B 8.60 9.833e-16 632-645
789	BL00740	MAM domain proteins.	BL00740B 19.76 5.378e-12 174-195 BL00740C 15.93 4.000e-11 684-695
793	PD00289	PROTEIN SH3 DOMAIN REPEAT PRESYNA.	PD00289 9.97 9.500e-12 102-116
795	BL00038	Myc-type, 'helix-loop-helix' dimerization domain proteins.	BL00038A 13.61 3.400e-09 66-82
800	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 5.050e-15 233-272
804	BL00663	Vinculin family talin- binding region proteins.	BL00663G 24.17 1.000e-40 364-414 BL00663K 21.52 9.816e-40 735-790 BL00663I 27.27 4.447e-35 514-568 BL00663J 18.16 3.000e-33 690-727 BL00663L 20.67 9.118e-27 802-838 BL00663F 20.78 2.000e-25 292-333 BL00663H 27.09 1.703e-24 436-489 BL00663C 22.59 2.853e-23 104-159 BL00663B 27.86 4.629e-23 42-96 BL00663D 24.77 3.789e-18 179-226 BL00663A 11.51 2.350e-15 18-39 BL00663E 21.19 9.566e-10 227-265
808	PR00010	TYPE II EGF-LIKE SIGNATURE	PR00010C 11.16 7.545e-10 3968-3979
809	PR00010	TYPE II EGF-LIKE SIGNATURE	PR00010C 11.16 7.545e-10 3882-3893

SEQ ID NO:	Accession No.	Description	Results*	
810	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 2.929e-10 163-196	
811	PR00205	CADHERIN SIGNATURE	PR00205B 11.39 9.182e-15 243-261 PR00205A 14.73 1.000e-12 168-184 PR00205C 13.65 1.783e-12 503-518 PR00205B 11.39 9.294e-11 463-481	
813	PR00456	RIBOSOMAL PROTEIN P2 SIGNATURE	PR00456E 3.06 5.146e-11 313-328 PR00456E 3.06 5.146e-11 314-329 PR00456E 3.06 5.146e-11 315-330 PR00456E 3.06 7.938e-10 312-327 PR00456E 3.06 7.938e-10 316-331	
818	BL01071	grpE protein.	BL01071A 24.88 8.277e-21 78-124 BL01071B 18.21 5.286e-15 195-219	
826	DM00813	AMINOTRANSFERASES CLASS-V PYRIDOXAL- PHOSPHATE ATTACHMENT SI.	DM00813A 20.30 8.898e-17 231-260	
828	BL00415	Synapsins proteins.	BL00415P 2.37 9.814e-09 242-278	
830	PF00632	HECT-domain (ubiquitin- transferase).	PF00632C 20.66 5.186e-23 1534-1566 PF00632B 18.45 8.393e-22 1480-1508	
831	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 9.695e-09 117-150	
832	PD01066	PROTEIN ZINC FINGER ZINC-FINGER METAL- BINDING NU.	PD01066 19.43 4.231e-33 12-51	
834	BL00120	Lipases, serine proteins.	BL00120B 11.37 5.846e-09 1319-1334	
836	DM00813	AMINOTRANSFERASES CLASS-V PYRIDOXAL- PHOSPHATE ATTACHMENT SI.	DM00813A 20.30 8.898e-17 38-67	
838	PD00289	PROTEIN SH3 DOMAIN REPEAT PRESYNA.	PD00289 9.97 8.000e-12 69-83	
840	BL00053	Ribosomal protein S8 proteins.	BL00053B 14.56 1.000e-08 900-918	
841	PR00970 .	ARGININE ADP- RIBOSYLTRANSFERASE SIGNATURE	PR00970D 9.96 3.357e-17 129-146 PR00970A 17.73 8.600e-17 30-52 PR00970E 11.23 6.464e-15 177-193 PR00970B 16.37 2.756e-11 58-77 PR00970C 11.05 9.357e-11 89-104	
842	BL00250	TGF-beta family proteins.	BL00250A 21.24 7.120e-25 114-150 BL00250B 27.37 4.774e-18 178-214	
846	BL00240	Receptor tyrosine kinase class III proteins.	BL00240B 24.70 7.488e-10 156-180	
848	BL01095	Chitinases family 18 proteins.	BL01095B 10.82 5.500e-14 24-36 BL01095C 10.76 7.207e-10 246-258	
849	BL00027	'Homeobox' domain proteins.	BL00027 26.43 2.500e-34 300-343	
850	PR00318	ALPHA G-PROTEIN (TRANSDUCIN) SIGNATURE	PR00318A 7.84 7.088e-09 188-204	
851	PF00969	Class II histocompatibility antigen, beta domain proteins.	PF00969A 22.07 5.846e-29 12-55 PF00969B 9.97 6.211e-25 56-92 PF00969C 27.72 7.324e-16 95-145	
852	BL00269	Mammalian defensins proteins.	BL00269B 19.17 6.824e-21 34-63 BL00269A 8.53 6.108e-18 1-21	
853	PF00777	Sialyltransferase family.	PF00777B 29.69 8.767e-10 407-450	
856	DM00191	w SPAC8A4.04C RESISTANCE SPAC8A4.05C DAUNORUBICIN.	DM00191D 13.94 9.083e-10 100-139	

SEQ ID NO:	No.		Results*
857	PR00823	PANCREATIC LIPASE	PR00823A 18.01 3.143e-14 19-37
		SIGNATURE	PR00823C 6.88 6.164e-12 56-69
859	BL00678	Trp-Asp (WD) repeat proteins proteins.	BL00678 9.67 6.684e-09 243-254
860	BL00028	Zinc finger, C2H2 type,	BL00028 16.07 8.650e-13 425-442
000	DECCOZO	domain proteins.	BL00028 16.07 5.696e-12 508-525
		domain proteins.	BL00028 16.07 8.826e-12 564-581
			BL00028 16.07 7.577e-11 201-218
			BL00028 16.07 7.577e-11 536-553
			BL00028 16.07 7.923e-11 341-358
			BL00028 16.07 8.615e-11 285-302
			BL00028 16.07 1.600e-10 592-609
			BL00028 16.07 2.200e-10 229-246
			BL00028 16.07 3.400e-10 257-274
			BL00028 16.07 6.100e-10 313-330
			BL00028 16.07 7.000e-10 369-386
		İ	BL00028 16.07 8.200e-10 397-414
			BL00028 16.07 5.114e-09 620-637
864	BL01126	Elongation factor Ts	BL01126A 18.48 5.011e-10 2637-2680
004	BEOTIZO	proteins.	
865	BL00353	HMG1/2 proteins.	BL00353B 11.47 1.330e-13 95-145
		•	BL00353B 11.47 5.692e-11 353-403
866	BL00972	Ubiquitin carboxyl-terminal	BL00972A 11.93 4.600e-18 173-191
		hydrolases family 2 proteins.	BL00972D 22.55 1.947e-13 576-601
			BL00972E 20.72 2.038e-11 618-640
867	BL00383	Tyrosine specific protein	BL00383E 10.35 2.756e-12 255-266
	}	phosphatases proteins.	
872	BL00030	Eukaryotic RNA-binding	BL00030B 7.03 5.737e-09 69-79
	1	region RNP-1 proteins.	
873	BL00303	S-100/ICaBP type calcium	BL00303B 26.15 4.405e-19 50-87
		binding protein.	BL00303A 21.77 8.765e-15 3-40
874	BL00523	Sulfatases proteins.	BL00523A 13.36 6.500e-17 41-58
			BL00523B 8.64 5.909e-15 89-101
			BL00523C 12.64 5.500e-13 140-151
			BL00523D 9.89 9.438e-11 293-305
877	BL00183	Ubiquitin-conjugating	BL00183 28.97 1.000e-40 42-90
001	DD 00001	enzymes proteins.	PR00081B 10.38 6.727e-11 116-128
881	PR00081	GLUCOSE/RIBITOL	PR00081B 10.38 6.727e-11 116-128
		DEHYDROGENASE	FRUUUBIA 10.33 3.1008-10 40-38
000	71.00100	FAMILY SIGNATURE	BL00183 28.97 1.391e-39 50-98
882	BL00183	Ubiquitin-conjugating	DLUUJ83 28.97 1.3916-39 30-98
		enzymes proteins.	DI 00102 20 07 1 201 20 50 02
883	BL00183	Ubiquitin-conjugating	BL00183 28.97 1.391e-39 50-98
		enzymes proteins.	77.00000 0 0 10 0 00 00 00 00 00 00 00 00 0
888	BL00027	'Homeobox' domain	BL00027 26.43 2.929e-30 232-275
		proteins.	

Results include Accession number, sub type, eMatrix p-value and the position of the signature.

TABLE 4

SEQ ID NO:	Pfam Model	Description	E-value	Pfam Score
45	Rap_GAP	Rap/ran-GAP	6.2 e -121	415.2
45 46	zf-C2H2	Zinc finger, C2H2 type	7.4e-65	228.9
	WD40	WD domain, G-beta repeat	0.00017	28.4
52	WD40	WD domain, G-beta repeat	1 3e-19	78.6
65	+	Carboxylesterases	2 le-128	440.0
83	COesterase	Hexokinase	0	2009.4
84	hexokinase		1e-135	464.2
85	zf-C2H2	Zinc finger, C2H2 type	3 2e-125	424.7
186	GTP EFTU	Elongation factor Tu family		955.3
187	myosin_head	Myosin head (motor domain)	1.5e-283	53.7
188	Glyco_transf_8	Glycosyl transferase family 8	4e-12	
189	tubulin	Tubulin/FtsZ family	3 2e-293	987.5
192	pkinase	Eukaryotic protein kinase domain	7 9e-85	295.2
193	SH3	SH3 domain	1.2e-18	75.4
197	Syntaxin	Syntaxin	0.074	-75.1
198	SCAN	SCAN domain	5.4e-67	236.0
199	F-box	F-box domain	0.0002	28.1
501	FHA	FHA domain	1.7e-13	58.3
502	Collagen	Collagen triple helix repeat (20 copies)	6.5e-197	667.6
507	PH	PH domain	3e-15	59.5
508	CH	Calponin homology (CH) domain	0.0069	16.3
512	ATPIG1 PLM_M	ATP1G1/PLM/MAT8 family	5.7e-31	116.3
312	AT8	,		
516	DnaJ	DnaJ domain	1.4e-24	95.1
519	DnaJ	DnaJ domain	6.8e-26	99.5
522	Glycos transf_2	Glycosyl transferases	1.2e-13	58.8
		Globin	4.1e-38	137.3
523	globin myosin head	Myosin head (motor domain)	0	1057.8
526		Acetyltransferase (GNAT) family	5e-11	50.1
529	Acetyltransf	MSP (Major sperm protein) domain	1.7e-16	68.2
530	MSP_domain	7 transmembrane receptor (Secretin	1.3e-59	211.5
531	7tm_2	family)	1.50-57	
534	Tropomyosin	Tropomyosin	7e-177	553.3
535	Tropomyosin	Tropomyosin	3.1c-173	541.9
538	LRR	Leucine Rich Repeat	2.9e-23	90.7
539	tRNA-synt_lb	tRNA synthetases class I (W and Y)	7.9e-79	275.3
540	PAS PAS	PAS domain	2.8e-05	24.9
	ferritin	Ferritin	9.9e-116	391.6
541		Cyclophilin type peptidyl-prolyl cis-tr	3.5e-33	117.6
546	pro isomerase	KH domain	0.0004	27.1
549	KH-domain	Glycosyl transferase family 8	0.0004	-47.7
551	Glyco_transf_8		2.6e-22	87.5
554	zf-C2H2	Zinc finger, C2H2 type		
555	Arylesterase	Arylesterase	2.3e-211	715.6
556	G-patch	G-patch domain	2.4e-17	71.1
557	DEAD	DEAD/DEAH box helicase	8.7e-67	214.2
558	Methyltransf_4	Putative methyltransferase	0.0095	-48.4
559	DSPc	Dual specificity phosphatase, catalytic dom	4.8e-70	246.1
563	IPPT	IPP transferase	6.7 e -66	232.4
566	zf-C2H2	Zinc finger, C2H2 type	2.6e-19	77.6
570	RNA_pol_L	RNA polymerases L / 13 to 16 kDa	0.043	-12.1
571	Armadillo seg	subunit Armadillo/beta-catenin-like repeat	8.6e-33	122.4
		WD domain, G-beta repeat	1.1e-65	231.6
574	WD40		1.2e-19	78.7
576	PAP2	PAP2 superfamily	3e-25	97.3
577	Defensin propep	Defensin propeptide		57.2
578	ig	Immunoglobulin domain	3.5e-16	
580	PI3 PI4_kinase	Phosphatidylinositol 3- and 4-kinase	6.5e-93	322.1

SEQ ID NO:	Pfam Model	Description	E-value	Pfam Score
585	GBP	Guanylate-binding protein, N-terminal domain	4.3e-165	548.2
586	PGM_PMM_I	Phosphoglucomutase/phosphomannom utase, alp	7.6e-06	4.4
587	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	1.3e-11	41.9
588	MMR HSR1	GTPase of unknown function	5.9e-48	172.7
590	zf-DHHC	DHHC zinc finger domain	1.8 e -36	134.6
591	Ribosomal_S3_C	Ribosomal protein S3, C-terminal domai	1.3e-07	28.0
592	LIM	LIM domain	4.4e-27	103.4
594	pkinase	Protein kinase domain	3.7e-77	269.7
596	PX	PX domain	2.2e-17	71.2
	Cadherin C term	Cadherin cytoplasmic region	3.3e-88	306.5
599	FHA	FHA domain	3.4e-20	80.5
600		Acetyltransferase (GNAT) family	3.2e-17	70.6
601	Acetyltransf	Nucleosome assembly protein (NAP)	5.5e-12	46.4
604	NAP_family	Nucleosome assembly protein (NAI)	1e-28	108.9
605	RhoGAP	RhoGAP domain	0.00022	28.0
606	Armadillo_seg	Armadillo/beta-catenin-like repeat	5.9e-77	269.1
607	pkinase	Protein kinase domain		
608	zf-C2H2	Zinc finger, C2H2 type	5.4e-110	378.8
609	ras	Ras family	1.2e-16	52.8
610	ank	Ank repeat	1 6e-08	41.8
612	pkinase	Protein kinase domain	1.6e-69	244.3
613	WD40	WD domain, G-beta repeat	4 7e-55	196.3
614	UBA	UBA/TS-N domain	3 6e-12	53.9
615	Zip	ZIP Zinc transporter	8 1e-59	208.8
618	sushi	Sushi domain (SCR repeat)	1 3e-58	208.2
619	K tetra	K+ channel tetramerisation domain	1 3e-19	78.6
	CAP GLY	CAP-Gly domain	1 9e-48	174.3
621	TCTP	Translationally controlled tumor protein	5.2e-109	375.5
622		Ubiquitin-conjugating enzyme	0.0046	-43.3
628 629	UQ_con DPPIV_N_term	Dipeptidyl peptidase IV (DPP IV) N- termi	5.1e-07	-82.1
630	DPPIV_N_term	Dipeptidyl peptidase IV (DPP IV) N- termi	5.5e-07	-83.2
631	efhand	EF hand	2.3e-14	61.1
632	Pribosyltran	Phosphoribosyl transferase domain	4.3e-37	136.7
635	ank	Ank repeat	1.8e-25	98.0
636	MHCK_EF2_kinas	MHCK/EF2 kinase domain family	1.2e-12	5.6
627	DUF221	Domain of unknown function DUF221	1.2e-89	311.2
637	bromodomain	Bromodomain	2.2e-29	106.0
	ubiquitin	Ubiquitin family	2.2e-21	81.9
641		RIO1/ZK632.3/MJ0444 family	1.1e-07	-14.9
644	RIOI	Nucleoside diphosphate kinase	1.1e-52	188.4
646	NDK -f.C2HC4	Zinc finger, C3HC4 type (RING finger)	9.4e-12	42.4
649	zf-C3HC4	ABC transporter	7.9e-84	291.9
651	ABC_tran		3e-30	113.9
654	CUB	CUB domain	2.6e-09	-35.3
655	MHCK_EF2_kinas	MHCK/EF2 kinase domain family		
659	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	1.3e-11	41.9
661	UvrD-helicase	UvrD/REP helicase	0.078	9.7
663	TFIIS	Transcription factor S-II (TFIIS)	2e-22	87.9
664	dsrm	Double-stranded RNA binding motif	4.3e-42	153.3
665	rrm	RNA recognition motif.	0.002	24.8
669	OTU	OTU-like cysteine protease	1e-19	78.9
671	TFIIS	Transcription factor S-II (TFIIS)	2e-22	87.9
672	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	1.5e-05	22.3
0/2	AMP-binding	AMP-binding enzyme	1.6e-86	300.9

SEQ ID NO:	Pfam Model	Description	E-value	Pfam Score
679	MSP domain	MSP (Major sperm protein) domain	5.4e-18	73.2
680	MSP domain	MSP (Major sperm protein) domain	5.5e-11	49.9
683	RNase PH	3' exoribonuclease family	3e-42	153.8
684	lactamase B	Metallo-beta-lactamase superfamily	0.088	-15.6
686	tRNA anti	OB-fold nucleic acid binding domain	0.031	20.9
690	NHL	NHL repeat	8.2e-18	72.6
691	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	6.1e-09	33.2
693	WD40	WD domain, G-beta repeat	0.025	21.2
694	WD40	WD domain, G-beta repeat	1.1e-23	92.1
696	Choline kinase	Choline/ethanolamine kinase	1.6e-51	184.6
697	zf-C2H2	Zinc finger, C2H2 type	3.4e-74	259.9
698	cadherin	Cadherin domain	2.2e-05	31.3
701	PRK	Phosphoribulokinase / Uridine kinase	1.1e-79	278.1
701	1	family		
702	DnaJ	DnaJ domain	5e-26	99.9
888	PAX	'Paired box' domain	1.1e-87	304.7

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PDB Chain Start End PSI- Verify PMF	Start End PSI- Verify AA AA BLAST score	End PSI- Verify AA BLAST score	PSI- Verify BLAST score	Verify	ļ	PMF score		SeqFold score	Coumpound	PDB annotation
					,				BUILD OTHER METERS	OXIIXOBEDITCI ASE POZ DOMAIN
1b8q A 324 401 1.30E-07 0.39 0.99	324 401 1.30E-07 0.39	401 1.30E-07 0.39	1.30E-07 0.39	0.39		66:0			NEUKONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	NNOS, NITRIC OXIDE SYNTHASE
1be9 A 355 401 0.00014 -0.32 0.65	355 401 0.00014 -0.32	401 0.00014 -0.32	0.00014 -0.32	-0.32		0.65			PSD-95, CHAIN: A, CRIPT; CHAIN: B;	PEPTIDE RECOGNITION PEPTIDE RECOGNITION, PROTEIN LOCALIZATION
1116 329 414 1.30E-05 0.58 0.48	414 1.30E-05 0.58	414 1.30E-05 0.58	1.30E-05 0.58	0.58		0.48			INTERLEUKIN 16; CHAIN: NULL;	CYTOKINE LCF, CYTOKINE, LYMPHOCYTE CHEMOATTRACTANT FACTOR, PDZ DOMAIN
1kwa A 329 404 1.10F-11 0.26 0.8	329 404 1.10F-11 0.26	404 1.10F-11 0.26	1.10E-11 0.26	0.26		0.8			HCASK/LIN-2 PROTEIN; CHAIN: A, B;	KINASE HCASK, GLGF REPEAT, DHR; PDZ DOMAIN, NEUREXIN, SYNDECAN, RECEPTOR CLUSTERING, KINASE
Iqau A 325 410 7.00E-11 0.7 1	325 410 7.00E-11	410 7.00E-11	7.00E-11		0.7	-			NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1- 130); CHAIN: A;	OXIIXOREDUCTASE BETA-FINGER
19av A 324 396 5.60E-10 0.2 0.81	324 396 5.60E-10 0.2	396 5.60E-10 0.2	5.60E-10 0.2	0.5		0.81			ALPIIA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXLDE SYNTHASE (RESIDUES 1-130); CHAIN: B;	MEMBRANE PROTEIN/OXIDOREDUCTASE BETA- FINGER, HETERODIMER
3pdz A 324 414 2.80E-05 0.64 0.74	324 414 2.80E-05 0.64	414 2.80E-05 0.64	2.80E-05 0.64	0.64		0.74			TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN: A;	HYDROLASE PDZ DOMAIN, HUMAN PHOSPHATASE, HPTPIE, PTP-BAS, SPECIFICITY 2 OF BINDING
lalh A 109 189 1.70E.30 0.24 1	109 189 1.70E-30	189 1.70E-30	1.70E-30		0.24 1	_			QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
lath A 137 248 1.70E-23 -0.03 0.11	137 248 1.70E-23 -0.03	248 1.70E-23 -0.03	1.70E-23 -0.03	-0.03	 	0.11			QGSR ZINC FINGER PEPTIDE: CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
Lath A 194 278 4.20E-29 80.51	194 278 4.20E.29	278 4.20F-29	4.20E-29		8	80.	.08	51	QGSR ZINC FINGER PEPTIDE: CHAIN: A: DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
1alh A 195 277 4.20E-29 0.16 1	195 277 4.20E-29	277 4.20E-29	4.20E-29		0.16 1				QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC

PDB ID		Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
1	F								OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	FINGER, DNA-BINDING PROTEIN
lalh	1 7	A	205	276	6.80E-28	-0.05			QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
lalh	1 -	\	224	303	8.50E-28	-0.09	-		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
lalh	<u></u>	<	252	331	1.50E-28	-0.18	0.99		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE, CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
lath	ļ	e e	280	359	1.20E-30	-0.24			QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE, CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
lalh	 	<	53	133	5.10E-20	0.1	0.02		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGENDNA) COMPLEX (ZINC FINGENDNA), ZINC FINGER, DNA-BINDING PROTEIN
lalh	 	4	81	189	4.20E-25	-0.35	0.21		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
x ae	 	O O	108	189	1.20E-49	0.35			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Jme y		O	136	220	1.70E-38	0.38	0.95		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
1mc y	 	O	164	276	1.00E-44	0.11	86.0		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)

SEQ NO:	PDB ID	Chain	Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
446	J mc	U	223	303	1.70E-47	0.15	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	Ime y	ပ	27	501	5.10E-21	0.08	-0.13		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	Jme y	ပ	278	360	1.00E-48			97.55	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	Ime y	U	279	360	1.00E-48	0.11	_		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	lme y	U	53	133	5.10E-37	0.19	0.75		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	Ime y	ပ	80	161	1.00E-48	0.53	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	lme y	ပ	83	190	4.20E-27	-0.13	0.36		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	Ime y	Ö		105	5.10E-12	0.39	0.19		DNA; CHAIN: A. B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
446	<u>:</u>	V	108	192	1.20E-20			62.34	TRANSCRIPTION FACTOR	COMPLEX (TRANSCRIPTION

PDB annotation	REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)						COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX
Coumpound	JIIA; CHAIN: A; SS RNA GENE; CHAIN: B, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA, CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TEIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE:
SeqFold score				124.38			
PMF score		0.53	0.05		0.94	0.99	0.86
Verify score		0.22	-0.07		0.37	-0.14	0.11
PSI- BLAST		1.20E-20	1.00E-14	1.70E-36	1.70E-36	I.40E-35	3.40E-30
End		189	129	281	257	347	681
Start		601	09	108	109	204	09
Chain ID		A .	⋖	⋖	∢	<	<u>ح</u>
PDB ID		143	ā	1116	1116	1116	1116
SEQ	Ö	446	446	446	446	446	446

PDB annotation	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN					COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR EFFMENT YY1 ZINC 2
Coumpound		TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A B.
SeqFold score							
PMF score		0.87	0.92	96.0	0.72	_	86.0
Verify score		-0.04	-0.12	0.01	0.01	0.09	-0.01
PSI- BLAST		8.50E-32	2.80E-32	1.70E-28	1.50E-30	2.80E-35	9.80E-36
End		229	248	220	276	276	331
Start		18	901	911	144	691	198
Chain ID		<	U	U	U	ပ	U
PDB ID		1116	lubd	lubd	lubd	lubd	lubd
SEQ ID		446	446	446	446	446	446

pound PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	S. ADENO-COMPLEX (TRANSCRIPTION VIRUS PS FREGULATION/DNA) YING-YANG 1; FRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX COMPLEX		Ý Z		ASSOCIATED VIRUS PS COMPLEX (TRANSCRIPTION ASSOCIATED VIRUS PS REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	C. ADENO. COMPLEX (TRANSCRIPTION
ld Coumpound		YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA: CHAIN: A, B;	YY I, CHAIN: C; ADENO- ASSOCIATED VRUS P5 INITIATOR ELEMENT D? CHAIN: A, B;	YY1; CHAIN: C; ADENO-
PMF SeqFold score				105.01			
Verify Plv score sco		0.08	-0.08		0.01	0.24 0.11	0.37
PSI- BLAST		5.10E-32	2.80E-31	9.80E-36	3.40E-34	5.10E-23	5.10E-30
End		303	359	360	359	133	161
Start		205	221	253	256	27	53
Chain ID		O	O	O	U .	O .	ပ
PDB ID		lubd	lubd	1 ubd	1ubd	1ubd	lubd
SEQ D		446	446	446	446	446	446

ē E	PDB TD	Chain ID	Start	End	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
	+			·					CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Jubd	 	ပ	88	189	1.40E-32	0.11	66:0		YYI, CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)
2adr	-		53	107	1.70E-09	0.04	-0.19		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR
2gli	 	₹	601	331	2.80E-42	0.05	0.21		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)
2gli		<	116	275	5.10E-29	0.04	0.11		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli		4	165	359	1.10E-41	0.11	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)
2gli		<	195	333	2.80E-42			99.39	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli	1	<	207	330	3.40E-30	0.46	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli		∢	231	358	3.40E-32	0.27	_		ZINC FINGER PROTEIN GLII; CHAIN: A: DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli		V	32	091	3.40E-27	0.48	0.4		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-

PDB annotation	(DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	T.ASTICITY.	ETON, SPECTRIN,	IDING, ACTIN-	NG PROTEIN, G DI IPLICATION.	MAIN	TEIN TWO	TRIN, ALPHA	COILED-COILS.	TEIN	OTEIN TWO	TRIN, ALPHA	REGION, 22	COILED-COILS,	TEIN TWO	TRIN ALPHA	REGION, 22	COILED-COILS,	OTEIN	OTEIN TWO	TRIN, ALPHA	KEGION, 2.2	COILED-COILS,	STRUCTURAL PROTEIN	HELIX	ATENI TOIDI E
PDB an	BINDING PROTEIN/DNA,	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GI ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GI ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	CYTOSKEL FTON EL ASTICITY	MEMBRANE SKELETON, SPECTRIN,	CALMODULIN-BINDING, ACTIN-	BINDING, 3 CAPPING PROTEIN,	REPEAT, 4 SH3 DOMAIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINNER REGION, 2 2 TANDEM 3-HELIX COILED-COILS.	STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINKER REGION, 22	TANDEM 3-HELIX COILED-COILS, STRINGTHEAT PROTEIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINKER REGION, 22	TANDEM 3-HELIX COILED-COILS,	STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO	REPEATS OF SPECTRIN, ALPHA	HELICAL LINKER REGION, 2.2	TANDEM 3-HELIX COILED-COILS,	MEMORIURAL PROTEINE	BUNDLE, ALPHA HELIX	CONTENT A CTUT IS DO CITED IN TRIBIDITE
Coumpound		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	AT BUA SECTION: CHAIN:	NULL;				ALPHA SPECTRIN; CHAIN:	A, B, C;			ALPHA SPECTRIN; CHAIN:	A, B, C;			AT BUY SBECTBIN: CHAIN	ALFRA SFECTION, CHAIN.	, c, t,			ALPHA SPECTRIN; CHAIN:	A, B, C;			A INTERIOR INTERIOR	SSOI FROIEIN; CHAIN: A;	
SeqFold score					-	٠												5/.5										
PMF score		0.93		8	60:0				0.23	-			0.03									0.33				000	67.0	
Verify		-0.1	0.31	80	0.78				-0.21				-0.17									-0.07	•			5,	-0.32	
PSI- BLAST		2.80E-45	8.50E-31	7, 000	6.80E-10				1.00E-17				\$.60E-05					1./0E-23				1.70E-23				0 0 0 0	0.00028	
End		278	247		139				142			_	201				3	797				259					171	
Start AA		83	88		39				2	l			-	,			ţ	3/			į	38				-	2	
Chain ID		×.	4			-			4	:				:				⋖				٧					<	
PDB ID	+	2gli	2gli		laj3				leun				Cun					Icun				1cun					1110	
SEQ ID	Ë	446	446		447				447				447	-				447				447					4	

PDB annotation	PROTEIN		CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CHAPTER CASE 11OB TRD DOMAIN	CHAFERONE HOF, IFR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN,	PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	CHAPERONE HOP, TPR-DOMAIN,	PEPTIDE-COMPLEX, HELICAL	REFEAT, BSC/0, 2 BSF/0, FROTEIN BINDING	-	RECEPTOR 1. PTS1-BP, PEROXIN-5,	PTSI PROTEIN-PEPTIDE COMPLEX,	TETRATRICOPEPTIDE REPEAT, TPR, 2	+	_	RECEPTOR 1, PISI-BF, PEROAIN-3,	TETRATRICOPERTIDE COMPLEX,	HELICAL REPEAT		TRANSCRIPTION INHIBITOR BETA-	METHYLTRANSFERASE METHYLTRANSFERASE,	CHEMOTAXIS RECEPTOR METHYLATION	METHYLTRANSFERASE GNMT, S-	ADENOSYL-L-METHIONINEN: GLYCINE METHYLTRANSFERASE	
Coumpound	A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	a Contract of the Contract of	TPRZA-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	TPR2A-DOMAIN OF HOP;	CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	TPRI-DOMAIN OF HOP;	CHAIN: A, B; HSC70-	PEPTIDE; CHAIN: C, D;	PEROXISOMAL TARGETING	SIGNAL I RECEPTOR;	CHAIN: A, B; PTS1-	CONTAINING PEPTIDE;	CHAIN: C, D;	PEROXISOMAL TARGETING	SIGNAL I RECEPTOR;	CHAIN: A, B; PISI-	CHAIN: C, D;		TRANSCRIPTIONAL REPRESSOR TUPI; CHAIN: A, B, C;	CHEMOTAXIS RECEPTOR METHYLTRANSFERASE	CHER; CHAIN: NULL;	GLYCINE N-	METHYLTRANSFERASE; CHAIN: A, B;	
SeqFold score					.		_																			
PMF		0.16	0.11		0.27	0		0.3			91.0					0.49					0.19	0.29		0.21		
Verify score		-0.31	-0.19		-0.26	-0.49		0.41			-0.37		_			0.12					0.01	-0.22		-0.32		
PSI- BLAST		0.0012	1.70E-19		5.10E-05	0.0014		1.20E-08			0.0084					4.20E-07	-				5.10E-38	0.0098		2.80E-06		
End		322	265		190	220		186			278					580					213	146		148		
Start		256	27		110	128		71			128					289					4	2		83		
Chain ID		A	<		∀	A		V			<					ď					<			<		
PDB ID		Idun	lquu		lelr	lei		lelw			1fch					1 Ch	_			L	lerj	laf7		lxva		
SEQ ID	Ö	447	447		450	450		450			450					450				L	452	458		458		

	PDB Chain ID ID		Start E	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
	18/1		×		0.0042	-0.40	60.00		NULL;	LIM DOMAIN CONTAINING LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN, ZINC 2 FINGER
∞ .	1589 A	351	7.1	718	5.10E-33	0.11	0.16		CLATHRIN HEAVY CHAIN; CHAIN: A;	CLATHRIN CLATHRIN, TRISKELION, COATED VESICLES, ENDOCYTOSIS, SELF- 2 ASSEMBLY, ALPHA-ALPHA SUPERHELIX
Icxx	4	851	8	879	8600.0	-0.39	0.04		CYSTEINE AND GLYCINE- RICH PROTEIN CRP2; CHAIN: A;	SIGNALING PROTEIN LIM DOMAIN CONTAINING PROTEINS, METAL- BINDING PROTEIN
lcrz	V 2	225	48	481	0.0012	0.31	0.77		TOLB PROTEIN; CHAIN: A;	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHABETA FOLD
lcrz	Y 2	341	46	499	0.007	0.51	0.15		TOLB PROTEIN; CHAIN: A;	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHABETA FOLD
lerj	V .	167	40	405	1.70E-48	0.31	0.28		TRANSCRIPTIONAL REPRESSOR TUP 1; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
lerj	∢	181	25	200	5.10E-72	9.0	92.0		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
lerj	4	220	5.5	551	6.80E-63	0.31	0.15		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
l erj	V	282	55	599	8.50E-64	0.36	0.41		TRANSCRIPTIONAL REPRESSOR TUPI; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
lerj	¥	330	9	639	1.70E-61	-0.03	0.01		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C:	TRANSCRIPTION INHIBITOR BETA- PROPELLER
Igot	<u>a</u>	129	49	499	1.40E-84			101.01	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
NO:	3pg m		16	251	1.50E-67	0.72	-		TRANSFERASE (PHOSPHORYL) PHOSPHOGLYCERATE MUTASE (E.C.2.7.5.3) DE- PHOSPHO ENZYME 3PGM 4	
473	1bor		27	70	3.40E-07	-0.13	0.04		TRANSCRIPTION FACTOR PML, CHAIN: NULL;	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION
473	Ichc		28	73	5.60E-12	0.11	96.0		VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	
473	1chc		29	76	1.70E-11	-0.45	99.0		VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	
473	1fbv	⋖	27	71	8.40E-14	-0.26	0.51		SIGNAL TRANSDUCTION PROTEIN CBL; CHAÎN: A; ZAP-70 PEPTIDE; CHAÎN: B; UBIQUITIN-CONJUGATÎNG ENZYME E12-18 KDA UBCH7; CHAÎN: C;	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
473	1fbv	₹	29	71	1.70E-11	-0.36	9.0		SIGNAL TRANSDUCTION PROTEIN CBL, CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
473	1825	v	28	73	5.60E-13	-0.15	0.71		CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)
473	1825	¥	29	08	3.40E-06	0.21	0.43		CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)
473	Irmd		20	112	2.80E-25	-0.05	0.93		RAGI; CHAIN: NULL;	DNA-BINDING PROTEIN V(D)) RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR

SEQ ID	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
Š										CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN
473	1ти		23	114	5.10E-21	-0.26	0.86		RAGI; CHAIN: NULL;	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN
473	1rmd		7	114	2.80E-25			54.37	RAGI; CHAIN: NULL;	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN
473	lubd	U	68	194	3.40E-11	-0.51	0.06		YYI, CHAIN: C: ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
473	Sznf		140	167	0.0042	-0.67	0.12		ZINC FINGER DNA BINDING DOMAIN ZINC-FINGER (ZFY- 6T) (NMR, 13 STRUCTURES) 5ZNF 3	
475	1av1	4		206	0.00056			61.09	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	LIPID TRANSPORT APO A-I; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION
475	Icun	<	13	219	0.00098			51.86	ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
475	c Ihm		164	204	0.0014	-0.45	0.01		DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB)	

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									(DNA-BINDING 1HME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) 1HME 4	
475	Ihsm		164	210	0.00084	-0.39	0.12		DNA-BINDING HIGH MOBILITY GROUP PROTEIN 1 (HMG1) BOX 2, COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	
475	Iquu	4	_	222	4.20E-08			55.34	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN
477	1bq0		108	182	1.20E-19			72.19	DNAJ; CHAIN: NULL;	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK
477	1 եզ		110	181	1.20E-19	=	-		DNAJ; CHAIN: NULL;	CHAPERONE HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK
477	lelr	<	9	129	1.20E-19	0.38	-0.09		TPR2A-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING
477	leíw	4	9	123	8.50E-24	0.42	0.05		TPR1-DOMAIN OF HOP; CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN BINDING
477	1fch	4	7	108	1.00E-18	0	-0.02		PEROXISOMAL TARGETING SIGNAL I RECEPTOR; CHAIN: A, B; PTSI- CONTAINING PEPTIDE; CHAIN: C, D;	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT
477	1hdj		107	180	1.20E-19	1.05	-		HUMAN HSP40; CHAIN: NULL;	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE
477	Ihdj		107	187	1.40E-30			80.39	HUMAN HSP40; CHAIN: NULL;	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE
477	1hdj		601	176	1.40E-30	0.98	-		HUMAN HSP40; CHAIN: NULL;	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE
479	1bjf	A	43	76	0.00013	-0.35	0.39		NEUROCALCIN DELTA;	CALCTUM-BINDING CALCTUM-

Start	Start AA		End AA	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
								CHAIN: A, B;	BINDING, MYRISTOYLATION, NEURONAL SPECIFIC GUANYLATE 2
41 83 0.00014	83		0.00014		-0.18	0.92		EPIDERMAL GROWTH FACTOR RECEPTOR PATHWAY CHAIN: 4:	CYCLASE ACTIVATION SIGNALING PROTEIN CALCIUM BINDING, SIGNALING DOMAIN, NPF BINDING EW RINDING 2 EF-HAND
41 83 0.00028	83		0.00028		-0.3	0.42		CALCIUM VECTOR PROTEIN; CHAIN: A;	EH DOMAIN, SIGNALING PROTEIN METAL BINDING PROTEIN CAVP; EF- HAND FAMILY, CALCIUM BINDING PROTEIN NMR
41 107 8.40E-05	107		8.40E-05		0.54	0.51		CALCIUM-BINDING PROTEIN CALMODULIN COMPLEXED WITH CALMODULIN-BINDING DOMAIN OF ICDM 3 CALMODULIN-BENDENT RAMODULIN-DEPENDENT PROTEIN KINASF II ICDM 4	
41 103 4.20E-05	103	 	4.20E-05		0.07	0.13		CALCIUM-BINDING PROTEIN CALMODULIN (VERTEBRATE) ICLL 3	
43 98 0.00017	86		0.00017	†	0.28	0.12		CARDIAC TROPONIN C; CHAIN: A;	STRUCTURAL PROTEIN HELIX-TURN- HELIX
41 103 0.00014	103	 	0.00014		60.0	0.27		CALMODULIN; CHAIN: A;	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER
43 84 5.60E-05	84		5.60E-05		0.14	0.21		CALCIUM-BINDING PROTEIN NCS-1; CHAIN: A;	METAL BINDING PROTEIN YEAST FREQUENIN EF-HAND, CALCIUM
42 96 0.00028	96		0.00028		-0.43	0.05		RECOVERIN; CHAIN: NULL;	CALCIUM-BINDING PROTEIN CALCIUM-MYRISTOYL SWITCH. CALCUIM-BINDING PROTEIN
22 143 1.70E-24	143		1.70E-24		0.11	0.63		PROMYELOCYTIC LEUKEMIA ZINC FINGER	GENE REGULATION POZ DOMAIN; PROTEIN-PROTEIN INTERACTION
								PROTEIN PLZF; CHAIN: A;	DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, PROMYELOCYTIC LEUKEMIA, GENE PROGII ATION
52 168 1.10E-05	168	<u> </u>	1.10E-05		-0.06	69:0		SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19;

							, 			
PDB annotation	SKPI, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE SKP2 F-BOX; SKP1, SKP1, SKP2, SKP2, F-BOX, LRR, LEUCINE-RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE				INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	VIRUS/VIRAL PROTEIN, RECEPTOR CD155, PVR, HUMAN POLIOVIRUS, ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR COMPLEX, VIRUS/VIRAL PROTEIN, RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE
Coumpound		CYCLIN A/CDK2- ASSOCIATED P19; CHAIN: A, C; CYCLIN A/CDK2- ASSOCIATED P45; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	OXIDOREDUCTASE(OXYGE N(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH 4.5) 1GOF 3	OXIDOREDUCTASE(OXYGE N(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH 4.5) IGOF 3	OXIDOREDUCTASE(OXYGE N(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH 4.5) 1GOF 3	HEMOLIN; CHAIN: A, B,	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	POLIOVIRUS RECEPTOR; CHAIN: R; VP1; CHAIN: 1; VP2; CHAIN: 2; VP3; CHAIN: 3; VP4; CHAIN: 4;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH
SeqFold score										
PMF		96.0	0.86	-0.08	0.22	-0.18	-0.17	0.21	-0.14	0.17
Verify score		0.48	0.26	0.22	0.31	0.07	0	-0.04	0.03	0.05
PSI- BLAST		4.20E-05	0.00042	1.40E-31	8.50E-13	2.80E-14	1.70E-29	1.40E-12	1.70E-14	5.10E-13
End		141	164	577	588	592	396	396	381	392
Start		52	22	318	346	362	66	303	96	303
Chain ID		æ	В				<	Q	ಜ	3
PDB		Ifsl	1fs2	lgof	1gof	1gof	16ih	lcvs	Idgi	lev2
SEQ		480	480	480	480	480	481	481	481	481

_	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	<u>e</u>	e	*	V V	BLAST	score	score	score		
									FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD
	1ev2	O	303	392	5.10E-13	-0.2	0.15		FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D;	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2;
									FIBROBLAST GROWTH	IMMUNOGLOBULIN (IG)LIKE
									CHAIN: E, F, G, H;	2 SUBGROUP WITHIN IG-LIKE
-+				,						DOMAINS, B-TREFOIL FOLD
	<u>اد</u>	ပ	303	396	1.40E-12	0.02	0.1		FIBROBLAST GROWTH FACTOR 1: CHAIN: A R:	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1: FGFR1:
									FIBROBLAST GROWTH	IMMUNOGLOBULIN (IG) LIKE
									FACTOR RECEPTOR 1;	DOMAINS BELONGING TO THE I-SET
									CHAIN: C, D;	2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD
\vdash	1 f 2q	<	192	386	1.70E-16	-0.2	0.04		HIGH AFFINITY	IMMUNE SYSTEM FC-EPSILON RI-
									IMMUNOGLOBULIN EPSII ON PECEPTOR CHAIN:	ALPHA; IMMUNOGLOBULIN FOLD, GI VCOPPOTEIN PECEPTOR 1GE.
									A;	BINDING 2 PROTEIN
-	lfhg	¥	289	381	3.40E-17	0.13	-0.07		TELOKIN, CHAIN: A	CONTRACTILE PROTEIN
										IMMUNOGLOBULIN FOLD, BETA BARREI
\leftarrow	Ikoa		292	382	8.50E-16	90.0	0.03		TWITCHIN; CHAIN: NULL;	KINASE KINASE, TWITCHIN,
-+-	1-36			, ,		200				INTRASTERIC REGULATION
\longrightarrow	2fcb	4	161	384	1.70E-14	-0.06	0.01		FC GAMMA RIIB: CHAIN: A;	IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM
-+										
	lol.	<	281	363	0.00014	-0.06	0.09		TPR2A-DOMAIN OF HOP: CHAIN: A; HSP90-PEPTIDE	CHAPERONE HOP, TPR-DOMAIN, PEPTIDE-COMPLEX, HELICAL
+									MEEV U, CHAIN, B,	NETERI, ASP 30, 2 FACIEIN BINDING
	la8s		152	296	0.0017	0.45	0.04		CHLOROPEROXIDASE F;	HALOPEROXIDASE
									CIONIN NOEE,	HALOPEROXIDASE,
-										OXIDOREDUCTASE, PROPIONATE COMPLEX
	1c4x	∢	178	331	0.00012	0.29	0.47		2-HYDROXY-6-OXO-6-	HYDROLASE BPHD, HYDROLASE,
\rightarrow									DIENOATE CHAIN: A;	reb Degrada IION
	1c7j	<	43	909	3.40E-93	0.39	-		PARA-NITROBENZYL FSTERASE: CHAIN: A:	HYDROLASE PNB ESTERASE; ALPHA- BETA HYDROI ASE DIBECTED
4									LOI LIVAGE, CHAINT. A,	DELA ILI DAVICASE, DINECLED

PDB annotation	EVOLUTION, ORGANIC ACTIVITY, 2 PNB ESTERASE	LIPASE ESTERASE, SUBSTRATE/PRODUCT-BOUND ICLE 9	LPASE ESTERASE, SUBSTRATE/PRODUCT-BOUND ICLE 9	HYDROLASE LINB, 1,3,4,6- TETRACHLORO-1,4- CYCLOHEXADIENE DEHALOGENASE, LINDANE, BIODEGRADATION, ALPHA/BETA-HYDROLASE	HYDROLASE (SERINE ESTERASE) HYDROLASE (SERINE ESTERASE), HYDROLASE, SERINE ESTERASE, 2 SYNAPSE, MEMBRANE, NERVE, MUSCLE, SIGNAL, NEUROTRANSMITTER 3 DEGRADATION, GLYCOPROTEIN, GPI-ANCHOR, ALTERNATIVE SPLICING	CHOLINESTERASE SERINE HYDROLASE, NEUROTRANSMITTER CLEAVAGE, CATALYTIC 2 TRIAD, ALPHA/BETA HYDROLASE	HYDROLASE ALPHA/BETA HYDROLASE FOLD	HYDROLASE ALPHA/BETA HYDROLASE FOLD	HYDROLASE BILE SALT ACTIVATED LIPASE, ESTERASE, CATALYTIC DOMAIN	SERINE HYDROLASE SERINE HYDROLASE, DEGRADATION OF BREFELDIN A, ALPHA/BETA 2 HYDROLASE FAMILY	
Coumpound		CHOLESTEROL ESTERASE; ICLE 4 CHAIN: A, B, ICLE 5	CHOLESTEROL ESTERASE; 1CLE 4 CHAIN: A, B; ICLE 5	HALOALKANE DEHALOGENASE; CHAIN: A;	ACETYLCHOLINESTERASE; CHAIN: A;	ACETYLCHOLINESTERASE; CHAIN: A;	SERINE HYDROLASE; CHAIN: A;	SERINE HYDROLASE; CHAIN: A;	BILE SALT ACTIVATED LIPASE; CHAIN: A;	BREFELDIN A ESTERASE; CHAIN: A, B;	HYDROLASE LIPASE (E.C.3.1.1.3) (TRIACYLGLYCEROL LIPASE) COMPLEXED WITH
SeqFold score		181.19				ı					179.92
PMF			_	0.57	-	-	0.58	0.42	-	0.1	
Verify score			0.21	0.6	0.61	0.59	0.17	-0.05	0.5	-0.05	
PSI- BLAST		1.20E-73	1.20E-73	0.0084	0	0	3.40E-28	5.60E-39	0	5.10E-20	1.20E-71
End		581	593	379	611	612	346	283	612	334	581
Start		42	89	163	04	40	142	73	44	83	42
Chain TO		4	Y	«	⋖	∢	<	¥	<	<	
PDB ID		Icle	lcle	lcv2	1dx4	lea5	levq	levq	1f6w	1 jkm	11pp
SEQ ID		483	483	483	483	483	483	483	483	483	483

SEQ	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
Ö									ILPP 3 HEXADECANESULFONATE ILPP 4 ILPP 71	
483	Прр		89	593	1.20E-71	0.22			HYDROLASE LIPASE (E C.3 1 1 3) (TRIACYLGLYCEROL LIPASE) COMPLEXED WITH 1LPP 3 HEXADECANESULFONATE 1LPP 4 1LPP 71	
483	Imaa	<	38	612	0			368.25	ACETYLCHOLINESTERASE; CHAIN: A, B, C, D;	HYDROLASE MACHE; HYDROLASE, SERINE ESTERASE, ACETYLCHOLINESTERASE, TETRAMER, 2 HYDROLASE FOLD, GLYCOSYLATED PROTEIN
483	Imaa	<	38	612	0	0.72			ACETYLCHOLINESTERASE, CHAIN: A, B, C, D;	HYDROLASE MACHE; HYDROLASE, SERINE ESTERASE, ACETYLCHOLINESTERASE, TETRAMER, 2 HYDROLASE FOLD, GLYCOSYLATED PROTEIN
483	lqe3	4	40	602	1.70E-89			242.59	PARA-NITROBENZYL ESTERASE; CHAIN: A;	HYDROLASE PNB ESTERASE, ALPHA- BETA HYDROLASE DIRECTED EVOLUTION
483	lqe3	∢	\$	599	1.70E-89	0.33			PARA-NITROBENZYL ESTERASE; CHAIN: A;	HYDROLASE PNB ESTERASE; ALPHA- BETA HYDROLASE DIRECTED EVOLUTION
483	1qfm	V	26	397	5.60E-57	0.15	0.11		PROLYL OLIGOPEPTIDASE; CHAIN: A;	HYDROLASE PROLYL ENDOPEPTIDASE, POST-PROLINE CLEAVING PROLYL OLIGOPEPTIDASE, AMNESIA, ALPHA/BETA-HYDROLASE, BETA- 2 PROPELLER
483	lqfm	<	87	350	1.20E-35	0	0.03		PROLYL OLIGOPEPTIDASE; CHAIN: A:	HYDROLASE PROLYL ENDOPEPTIDASE, POST-PROLINE CLEAVING PROLYL OLIGOPEPTIDASE, AMNESIA, ALPHA/BETA-HYDROLASE, BETA-2 PROPELLER
483	lthg		46	280	5.10E-80			210.1	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E C 3 1 1.3)	

PDB annotation			HYDROLASE BILE SALT ACTIVALED LIPASE, BILE SALT STIMULATED HYDROLASE, SERINE ESTERASE, LIPASE	HYDROLASE BILE SALT ACTIVALED LIPASE, BILE SALT STIMULATED HYDROLASE, SERINE ESTERASE, LIPASE	HEXOKINASE ATP/.D-HEXOSE-6- PHOSPHOTRANSFERASE; HEXOKINASE, PHOSPHOTRANSFERASE	HEXOKINASE ATP/:D-HEXOSE-6- PHOSPHOTRANSFERASE; HEXOKINASE. PHOSPHOTRANSFERASE	TRANSFERASE STRUCTURALL.Y HOMOLOGOUS DOMAINS	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA
Coumpound	TRIACYLGLYCEROL HYDROLASE 1THG 3	HYDROLASE(CARBOXYLIC ESTERASE) LIPASE (E.C.3.1.1.3) TRIACYLGLYCEROL HYDROLASE ITHG 3	CHOLESTEROL ESTERASE; CHAIN: NULL;	CHOLESTEROL ESTERASE; CHAIN: NULL;	HEXOKINASE; CHAIN: A, B;	HEXOKINASE, CHAIN: A, B,	HEXOKINASE TYPE I; CHAIN: N;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE, CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	
SeqFold score			318.91			179.92					
PMF		_		-	-		_	0.11	0.3	0.94	0.11
Verify		0.49		0.54	1.19		1.21	-0.49	-0.36	0.16	-0.34
PSI- BLAST		5.10E-80	0	0	0	0	0	1.70E-18	1.70E-23	6.80E-24	1.70E-30
End		583	618	612	910	116	913	204	232	260	204
Start		47	39	44	ļ-	_	91	127	152	180	126
Chain ID					A	<	z	«	<	<	C
PDB ID		Ithg	2bce	2bce	1bg3	1bg3	lcza	lalh	laih	lalh	lme y
SEQ ID		483	483	483	484	484	484	485	485	485	485

	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
	1								PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
lme y	9	O	151	232	5.10E-38	-0.18	0.69		DNA; CHAM: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Ime y	ဥ	O	621	260	8.50E-41	-0.01	0.99		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
= >	J me	O	207	288	1.20E-42	0.05			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER. PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
= >	1me y	O	235	316	1.40E-43	0.27			DNÁ; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
- ×	Ime y	ى ن	291	372	2.80E-47	0.25	_		DNA; CHAIN; A. B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
- ×	lme y	O	291	372	5.10E-47	0.25	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
- >	Ime y	U	319	400	1.50E-48	0.67			DNA, CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN, CHAIN: C, F, G;	COMPLEX (ZINC FINGEN/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
->	1me y	U	319	400	2.80E-51	0.67			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2

SEQ ID NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
										CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
485	lme y	Ü	347	428	5.10E-49	0.48	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA
									PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
										CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
485	Ime	S	375	456	3.40E-49	69.0	-		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
	>								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
									PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
										(ZINC FINGER/DNA)
485	me >	ပ _	375	426	4.20E-50	69.0	<u></u>		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC ENGEP PROTEIN DNA
	`								PROTEIN: CHAIN: C. F. G.	INTERACTION, PROTEIN DESIGN, 2
										CRYSTAL STRUCTURE, COMPLEX
										(ZINC FINGER/DNA)
485	Ime	ပ	375	457	4.20E-50			113.26	DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
	>				_				CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
									PROJEIN; CHAIN: C, F, G;	CRIVERACTION, PROTEIN DESIGN, 2
		_								CALISTAE STRUCTORE, COMPLEX
485	lme	O	403	484	1.00E-49	0.41	_		DNA; CHAIN: A. B. D. E.	COMPLEX (ZINC FINGER/DNA) ZINC
	×								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
_			_						PROTEIN; CHAIN: C, F, G,	INTERACTION, PROTEIN DESIGN, 2
										CRYSTAL STRUCTURE, COMPLEX
100	1		12.5	9:5	100	-0.0				(ZINC FINGER/DNA)
	e ,	ر	431	512	1.70E-50	0.02			DNA; CHAIN: A. B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
_	`								DEOTERN CHARN DE C.	FINGER, FROJEIN-DNA
									FROIEIN; CHAIN: C, F, G;	Chystal strictime Country
_										CRISIAL SIRUCIORE, COMPLEX
485	Ime	C	459	240	1 50F-50	0.02	-		DNA: CHAIN: A B D E	COMPLEY (ZING ENGER DNIA) ZING
_)		?	1.300-30	7.05	-		CONSTRICTS AND ENGTR	COMPLEA (ZINC FINGER/DINA) ZINC
									DDOTEIN: CHAIN: C E G.	FINGER, FROI EIN-DIA
_									() (1 () () () () () () () () () () () () ()	CRYSTAL STRIICTHE COMPLEX
\neg										(ZINC FINGENDNA)
485	l'inc	၁	487	899	8.50E-51	0.1	_		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
	>-								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
									PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
1										CRISIAL SIRUCIUNE, CUMPLEA

Coumpound PDB annotation	(ZINC FINGER/DNA)	CONSENSUS ZINC FINGER, PROTEIN-DNA) ZINC FINGER, PROTEIN-DNA DESIGN, 2 PROTEIN: CHAIN: C, F, G, INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)		CONSENSUS ZINC FINGER FINGER, PROTEIN-DNA	FROIEIN, CHAIN: C, r, C, (RYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	-	CONSENSUS ZINC FINGER FINGER, PROTEIN-DNA			CONSENSUS ZINC FINGER FINGER, PROTEIN-DNA	FROJEJIN, CHALIN. C, F, G; INTERACTION, FROJEJIN DESIGN, Z	(ZINC FINGER/DNA)	DNA: CHAIN: A. B, D, E, COMPLEX (ZINC FINGER/DNA) ZINC	ER	PROTEIN; CHAIN: C, F, G. INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX		CONSENSIS ZINC FINGER FINGER PROTEIN-DNA			TFIIIA; CHAIN: A, D; SS COMPLEX (TRANSCRIPTION RIBOSOMAL RNA GENE; REGULATION/DNA) COMPLEX		REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION INTRATION ZINC FINGER PROTEIN	TEILIA CHAIN A D. 5S COMPLEX (TRANSCRIPTION	_	RIBOSOMAL RNA GENE; REGULATION/DNA) COMPLEX
	dalla.	CONSENSU PROTEIN; C		DNA; CHAI	CONSENSU	rroiem, c	-	DNA; CHAI	CONSENSO		DNA; CHAI	CONSENSO	FROJEJIN,		DNA: CHAI	CONSENSU	PROTEIN; C		DAIA. CHA	CONSENSU	PROTEIN, C		TFIIIA; CHA	CHAIN: B, C, E, F;			TFIIIA CHA		KIBOSOMA
SeqFold																			1				_				-		_
PMF score	-	<u>-</u>		-				_			-	_			_				07.0			 ,	0.76				0.81	:	_
Verify score		0.07		0.03				0.28			0.15				0.36				0.70	0.50			-0.28				200	!	_
PSI- BLAST	1 605 50	1.30E-30		3.40E-50				1.70E-50			3.40E-50				1.00E-32				\$ 10E 07	3.10E-07			3.40E-27				3 40F-31		
End	703	960		624				652	_		089				683				176	0/1	_	 - ;	274		_		797	1	
Start	313	C) C		543				571			599				627				0,1	6+1			130				157	!	_
Chain ID		ر		၁				၁			C				S				C	5			⋖					:	
PDB ID	1	y Be		Ime	>		_	Imc	>		lme	<u>`</u>			Ime	`) 	`		146				1166	,	
SEQ NO:	307	483	_	485				485			485				485				707	-			485				485	:	_

PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III. 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE 111, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1;
Coumpound		TFIIIA; CHAIN: A. D: 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D: 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS
SeqFold score			120.46					
PMF		_		1	0.99	66.0	0.12	0.72
Verify score		0.07		0.11	0.01	0.1	-0.22	-0.27
PSI- BLAST		6.80E-34	5.10E-37	5.10E-37	3.40E-37	1.40E-36	5.10E-25	8.50E-27
End		353	484	521	633	682	232	260
Start		208	319	376	488	544	127	159
Chain ID		₹	A	<	4	<	U	O
PDB ID		116	1tf6	1116	11166	1116	lubd	1ubd
SEQ ID	ë Z	485	485	485	485	485	485	485

SEQ	PDB	Chain	Start	End	PSI.	Verify	PMF	SeqFold	Coumpound	PDB annotation
Εģ	8	e	VV VV	VΥ	BLAST	score	score	score		
									INITIATOR ELEMENT DNA; CHAIN: A, B;	TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	pqn1	O	182	288	3.40E-29	-0.04	98.0		YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	pqnı	ပ	215	316	1.20E-29	0.32	0.93		YYI, CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	pqnl	U	238	344	8.50E-32	0.18			YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	1 ubd	U	240	345	4.20E-46	0.02	96.0		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1, TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	lubd	U	268	372	7.00E-52	0.14	_		YY I, CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	lubd	C	296	400	9.80E-59	0.63			YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION

<u> </u>	5-YANG 1; 1ON, 71, ZINC 2 ROTEIN EX ATION/DNA)	ION 5-YANG 1; TION, 71, ZINC 2 ROTEIN EX ATION/DNA)	ION G-YANG 1; FION, Y1, ZINC 2 ROTEIN EX.	TON G-YANG 1; TION, Y1, ZINC 2 *ROTEIN EX.	TON G-YANG 1; TION, Y1, ZINC 2 PROTEIN .EX	JON G-YANG I; TION, YI, ZINC 2 PROTEIN .EX
PDB annotation	REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY I, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A. B:
SeqFold score			100.22			
PMF		-		-	-	-
Verify		0.04		0.26	0.1	0.09
PSI- BLAST		2.80E-59	2.80E-59	2.80E-55	8.40E-56	4.20E-59
End		456	457	512	296	652
Start		346	349	401	485	541
Chain ID		U	U	U	υ	ပ
PDB ID		lubd	1ubd	Iubd	lubd	lubd
SEQ ID	Ö	485	485	485	485	485

SEQ ID NO:	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
485	lubd	၁	579	089	3.40E-34	90.0	0.98		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
485	2gli	4	131	262	1.00E-26	-0.05	60.0		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	∢	179	346	5.60E-44	0.01	0.47		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	4	207	343	3.40E-31	0.4	_		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	<	263	402	1.40E-68	0.53	_		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	<	319	458	2.80E-77			110.36	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	<	319	458	2.80E-77	0.54	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	4	375	542	1.40E-73	0.01	0.88		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2 g li	۲	487	654	1.10E-73	-0.12	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEINDNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	K	516	681	1.40E-72	60.0	86.0		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C,	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI,

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
NO.	a	a	VV	AA	BLASI	score	score	score		
									D;	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	٧	523	651	1.40E-34	0.23	0.94		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
485	2gli	Ą	1551	679	3.40E-33	0.03	0.95		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI. ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
486	1d2e	Ą	58	454	0			627.43	ELONGATION FACTOR TU (EF-TU); CHAIN: A, B, C, D	RNA BINDING PROTEIN G-PROTEIN, BETA-BARREL
486	1d2e	A	58	454	0	1.12	_		ELONGATION FACTOR TU (EF-TU); CHAIN: A, B, C, D	RNA BINDING PROTEIN G-PROTEIN, BETA-BARREL
486	ldar		49	192	3.40E-17	-0.05	0.19		ELONGATION FACTOR G; CHAIN: NULL;	TRANSLATIONAL GTPASE EF-G RIBOSOMAL TRANSLOCASE, TRANSLATIONAL GTPASE
487	laca		1926	1954	0.0014	-0.46	0.11		ACYL-COENZYME A	
						_			BINDING PROTEIN ACYL-	
									PROTEIN (ACBP) COMPLEX	
	_					_			WITH IACA 3 PALMITOYL-	
						_			COENZYME A (NMR, 20 STRUCTURES) 1ACA 4	
487	157t	<	_	794	0			510.37	MYOSIN HEAVY CHAIN;	MYOSIN MYOSIN MOTOR
									CHAIN: A; MYOSIN REGIT ATORY LIGHT	
									CHAIN; CHAIN: Y; MYOSIN	
					-	 ,-			ESSENTIAL LIGHT CHAIN;	
487	1291	4	5	807	0	0.41	_		MYOSIN HEAVY CHAIN:	MYOSIN MYOSIN MOTOR
						-			CHAIN: A; MYOSIN	
						_			REGULATORY LIGHT	
									CHAIN; CHAIN: Y; MYOSIN	
									ESSENTIAL LIGHT CHAIN; CHAIN: Z:	
487	1br1	A		765	0			524.14	MYOSIN, CHAIN: A, B, C, D,	MUSCLE PROTEIN MDE, MUSCLE
487	17.1	٧	~	760		030	-		E, F, G, H;	PROTEIN
	101			/0/		0.08	-		MYOSIN, CHAIN: A, B, C, D,	MUSCLE PROTEIN MDE, MUSCLE

PDB ID		Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
	1								E, F, G, H;	PROTEIN
1br2	1	A	=	725	0	0.58	_		MYOSIN; CHAIN: A, B, C, D, E, F;	MUSCLE PROTEIN MUSCLE PROTEIN
1br2		A	=	739	0			470.06	MYOSIN; CHAIN: A, B, C, D, E, F;	MUSCLE PROTEIN MUSCLE PROTEIN
1btk		4	1394	1495	9.80E-08	-0.2	0.11		BRUTON'S TYROSINE KINASE; CHAIN: A, B,	TRANSFERASE BRUTON'S AGAMMAGLOBULINEMIA TYROSINE KINASE, BTK; TRANSFERASE, PH DOMAIN, BTK MOTIF, ZINC BINDING, X-LINKED 2 AGAMMAGLOBULINEMIA, TYROSINE-PROTEIN KINASE
1btn	 		1216	1306	8.40E-13	0.42	0.28		BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	SIGNAL TRANSDUCTION PROTEIN
l btn			1318	1382	0.00042	-0.39	0.07		BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5	SIGNAL TRANSDUCTION PROTEIN
lci.			796	933	2.80E-19	0.21	-0.19		COLICIN IA; CHAIN: NULL;	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN
loii	 		810	616	2.80E-18	0.03	-0.19	,	COLICIN IA, CHAIN: NULL,	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN
Icun	 	V V	726	940	1.40E-13	0.07	-0.14		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
Jcm	 	A	808	946	1.10E-14	0.27	-0.17		ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
1dfk		V	8	807	0	0.23	_		MYOSIN HEAD, CHAIN: A; MYOSIN HEAD, CHAIN: Y; MYOSIN HEAD, CHAIN: Z;	CONTRACTILE PROTEIN MYOSIN MOTOR, CONFORMATIONAL CHANGES
<u>=</u>	-	¥	1710	2002	4.20E-27	0.13	66:0		MOESIN; CHAIN: A, B; MOESIN; CHAIN: C, D;	MEMBRANE PROTEIN CRYSTAL STRUCTURE, MEMBRANE, FERM DOMAIN, TAIL DOMAIN
lez3	\vdash	٧	808	885	1.40E-11	0.37	-0.2		SYNTAXIN-1A; CHAIN: A. B,	ENDOCYTOSIS/EXOCYTOSIS

Chain		Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									TRANSDUCTION PROTEIN, ADAPTOR PROTEIN
A		1215	1308	2.80E-16	0.21	0.31		GRP1; CHAIN: A;	SIGNALING PROTEIN ARFI GUANINE NUCLEOTIDE EXCHANGE FACTOR AND PH DOMAIN
V		1710	2044	2.80E-26	0.08	-		RADIXIN; CHAIN: A;	CELL ADHESION 3 SUBDOMAINS, CYTOSKELETON, CELL ADHESION
	1,4	2	739	0			540.6	MYOSIN: CHAIN: NULL;	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELJUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL
	1	\$	725	0	9.0	-		MYOSIN; CHAIN: NULL;	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL
	1	_	671	0			479.56	MYOSIN; CHAIN: NULL;	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN
	i	2	129	0	0.63			MYOSIN, CHAIN: NULL;	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN
		1214	1315	2.80E-17	0.11	0.46		PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT 1PLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS 1PLS 4 (INS(G105-LEHHHHHH)) (NMR, 25 STRUCTURES) 1PLS 5	
		1342	1381	5.60E-05	-0.55	0.04		PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3	

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ΞÖ	3	3	¥ ¥	¥ ¥	DLASI	SCULE	score:	30016		
									WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS 1PLS 4 (INS(G105- LEHHHHHH)) (NMR, 25 STRUCTURES) 1PLS 5	
487	slq1		1394	1495	9.80E-12	0.02	-0.01		PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN HOMOLOGY DOMAIN) MUTANT IPLS 3 WITH LEU GLU (HIS)6 ADDED TO THE C TERMINUS IPLS 4 (INS(G105-LEHHHHHH)) (NMR, 25 STRUCTURES) IPLS 5	
487	1 pms		1211	1308	8.40E-15	10.0	-0.01		SOS 1; CHAIN: NULL;	SIGNAL TRANSDUCTION SON OF SEVENLESS; PLECKSTRIN, SON OF SEVENLESS, SIGNAL TRANSDUCTION
487	Ipms		1331	1381	0.00014	-0.14	0.05		SOS 1; CHAIN: NULL;	SIGNAL TRANSDUCTION SON OF SEVENLESS; PLECKSTRIN, SON OF SEVENLESS, SIGNAL TRANSDUCTION
487	lqqg	4	1331	1464	7.00E-07	-0.27	0.25		INSULIN RECEPTOR SUBSTRATE 1: CHAIN: A, B;	SIGNAL TRANSDUCTION IRS-1; BETA-SANDWHICH, SIGNAL TRANSDUCTION
487	lquu	4	767	973	1.30E-20	0.05	-0.13		HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN
487	2mys	<	2	801	0			419.33	MYÖSIN; CHAIN: A, B, C;	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN
487	2mys	<	4	775	0	0.53	г		MYOSIN; CHAIN: A, B, C;	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN
489	Itub	<	-	440	0			727.18	TUBULIN; CHAIN: A, B;	MICROTUBULES MICROTUBULES, ALPHA-TUBULIN, BETA-TUBULIN, GTPASE HELIX
489	ltub	<	-	440	0 .	8.0	_		TUBULIN; CHAIN: A, B;	MICROTUBULES MICROTUBULES, ALPHA-TUBULIN, BETA-TUBULIN, GTPASE HELIX
492	1a06		16	312	1.50E-87	0.35			CALCIUM/CALMODULIN-	KINASE KINASE, SIGNAL

PDB annotation	TRANSDUCTION, CALCIUM/CALMODULIN	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	TRANSFERASE TRANSFERASE, SERINE/THREONINE-PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE		PROTEIN KINASE CDK2, PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION,
Coumpound	DEPENDENT PROTEIN KINASE; CHAIN: NULL;	ż		TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-AMP\$- DEPENDENT PROTEIN (KINASE (E.C.2.1.137) (\$C/ARLYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHBITOR PKI(\$-24) AND 1THE DETERGENT MEGA-8 1APM 6 TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 IAPM 4 REPLACED BY ALA (\$S1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(\$-24) AND THE DETERGENT MEGA-8 IAPM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;
SeqFold score		121.63	90.87	158.35	120.73
PMF score					
Verify score				0.41	
PSI- BLAST		1.50E-87	1.70E-43	0	1.00E-57
End		318	316	315	314
Start		17	2		22
Chain ID				ш	
PDB UD		1a06	1a60	m lap	laqi
SEQ		492	492	492	492

PDB annotation	MITOSIS, INHIBITION	COMPLEX (KINASE/INHIBITOR) CDK6; PI9INK4D; CYCLIN DEPENDENT KINASE, CYCLIN DEPENDENT KINASE INHIBITORY 2 PROTEIN, CDK, INK4, CELL CYCLE. COMPLEX (KINASE/INHIBITOR)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	TRANSFERASE CSK; PROTEIN KINASE, C-TERMINAL SRC KINASE, PHOSPHORYLATION, 2 STAUROSPORINE, TRANSFERASE	PHOSPHOTRANSFERASE PROTEIN KINASE ICKI 18				
Coumpound	2	CYCLIN-DEPENDENT CYCLIN-DEPENDENT CYCLIN-DEPENDENT IXINASE INHIBITOR: CHAIN: B. D:	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P191NK4D; CHAIN: B; F	C-TERMINAL SRC KINASE; 1 CHAIN: A; F	CASEIN KINASE I DELTA; ICKI 6 CHAIN: A, B; ICKI 7	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) ICTP 3 (CATALYTIC SUBUNIT) ICTP 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK)
SeqFold score		117.12	139.32	120.46	82.9		156.27		152.26
PMF									
Verify score						0.42		0.32	
PSI- BLAST		2.80E-54	1.40E-59	1.40E-39	9.80E-51		0 .	0	0
End	T	303	308	286	303	315	333	315	330
Start		23	81	18	17	-	٤.		E .
Chain ID		4	V V	∀	V	ய	ല	ш	ய
PDB ID	-	1bi8	1blx	1byg	1cki	r k	lcm k	lctp	lctp
SEQ ID		492	492	492	492	492	492	492	492

PDB annotation			TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER		TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT,	номорімея	-		FIBROBLAST GROWTH FACTOR	TYROSINE-PROTEIN KINASE, ATP-	BINDING, 2 PHOSPHORYLATION,	\dashv		FIBROBLASI GROWIH FACTOR	TYROSINE-PROTEIN KINASE, ATP-	BINDING, 2 PHOSPHORYLATION,	RECEPTOR, PHOSPHOTRANSFERASE	PROTEIN KINASE CDK2;	TRANSFERASE, SERINE/THREONINE	PROTEIN KINASE, AIP-BINDING, 2	MITOSIS PHOSPHORYI.ATION	SERINE/THREONINE-PROTEIN	KINASE CSBP, RK, P38; PROTEIN	SER/THR-KINASE,	SERINE/THREONINE-PROTEIN	+	(TRANSFERASE/SUBSTRATE)	TYROSINE KINASE, SIGNAL	TRANSDUCTION,	KINASE/PEPTIDE SUBSTRATE/ATP
Coumpound		SUBUNIT) 1CTP 4	SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINE/THREONINE- PROTEIN K NASE PAK.	ALPHA; CHAIN: C, D;	SERINE/THREONINE-	ALPHA; CHAIN: A, B;	PROTEIN KINASE PAK- ALPHA; CHAIN: C. D;	FGF RECEPTOR 1; CHAIN: A.	B;				FGF RECEPTOR 1; CHAIN: A,	_ B;				HUMAN CYCLIN-	DEPENDENT KINASE 2;	CHAIN: NULL;		P38 MAP KINASE: CHAIN:	NULL;			a control of a con	INSULIN KECEPTOK; CHAIN: A PEPTIDE SHBSTRATE	CHAIN: B;		
SeqFold	31036	,						123.75					127.9					141.29				104 86	204:00			,	105.26			
PMF	31016		_																				,							-
Verify	31016		0.38		0.21																									
PSI-	DLASI		2.80E-69		5.10E-69			2.80E-38					1.20E-40					1.40E-60				2 ROF-45	7.00.7		•		9.80E-40			
End	¥		303		293			286					285					314				346	2		_		7.67		· · · -	
Start	*		23		4			=					12					22				~	<u> </u>			,	۷_			
Chain	9		O		၁			4					В														<	_		
PDB	<u> </u>		113m		113m			1 fgk)				1 fgk					l hcl				lian	ē				Ξ.			
SEQ	ΞÖ		492		492			492					492					492				492	764				492			

		SE,					H H	ė Z				LP 2,
PDB annotation	ANALOG), ENZYME, 3 COMPLEX (TRANSFERASE/SUBSTRATE)	TRANSFERASE INK3, TRANSFERASE, INK3 MAP KINASE, SERINETHREONINE PROTEIN 2 KINASE	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE; TRANSFERASE, MAP KINASE, SERINE/THREONINE-PROTEIN KINASE, 2 P38	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE- PROTEIN, 2 KINASE, ATP-BINDING, CALMODULIN-BINDING	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM; TRANSFERASE, SERINE/THREONINE- PROTEIN. 2 KINASE, A TP-BINDING. CALMODULIN-BINDING	TRANSFERASE MAP KINASE. SERINE/THREONINE PROTEIN KINASE, TRANSFERASE	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, SERINE/THREONINE-PROTEIN KINASE, MAP KINASE, 2 ERK2
	ANALOG), (TRANSFEI	TRANSFERASE JNK JNK3 MAP KINASE, SERINE/THREONIN KINASE	KINASE KII INTRASTEI	KINASE KII INTRASTER	KINASE KII INTRASTEI	TRANSFERASI ACTIVATED PI TRANSFERASI SERINE/THREC KINASE, 2 P38	KINASE RA PHOSPHOR GLYCOGEN TRANSFER PROTEIN, 2	KINASE RA PHOSPHOR GLYCOGEN TRANSFER PROTEIN. 2	TRANSFER SERINE/TH KINASE, TF	SERINE KIN	SERINE KIN TITIN, MUS	TRANSFERASE MITOC ACTIVATED PROTEIN ERK2; TRANSFERASE, SERINETHREONINE-F
Coumpound		C-JUN N-TERMINAL KINASE; CHAIN: NULL;	TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	ERK2; CHAIN: NULL;	TITIN; CHAIN: A, B;	TITIN; CHAIN: A, B;	EXTRACELLULAR REGULATED KINASE 2; CHAIN: NULL;
SeqFold		127.36		139.94		121.52	109.51		118.65	126.69		130.2
PMF score			_		_		_	_			_	
Verify score			0.27		0.4			0.6			0.49	
PSI- BLAST		7.00E-54	1.70E-70	3.40E-71	3.40E-71	1.40E-56	1.20E-81	1.20E-81	5.60E-50	4.20E-65	4.20E-65	4.20E-56
End		357	334	351	284	350	282	279	341	334	274	346
Start AA		∞	22		17	4	22	23	18	61	22	=
Chain ID				4	V					٧	∢	
PDB ID		ljnk	Ikoa	Ikob	lkob	1p38	1 phk	l phk	lpm c	1tki	11ki	3erk
SEQ ID NO:		492	492	492	492	492	492	492	492	492	492	492

PDB annotation	LIPID TRANSPORT APO A-1; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	TRANSFERASE IL-2-INDUCIBLE T- CELL KINASE; TRANSFERASE, REGULATORY INTRAMOLECULAR COMPLEX, KINASE	COMPLEX (ADAPTOR PROTEIN/PEPTIDE) ASH, GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2; COMPLEX (ADAPTOR PROTEIN/PEPTIDE), SH3 DOMAIN, 2 GUANINE-NUCLEOTIDE RELEASING FACTOR	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE), SIGNAL TRANSDUCTION, 2 SH3 DOMAIN			ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	MEMBRANE PROTEIN FOUR HELIX BUNDLE, ALPHA HELIX	TRANSFERASE PROTO-ONCOGENE TYROSINE KINASE; PROTO- ONCOGENE, TRANSFERASE,
Coumpound	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	ITK; CHAIN: NULL;	GRB2; CHAIN: A; SOS; CHAIN: B;	ABL TYROSINE KINASE; CHAIN: A, C, E, G; PEPTIDE P41; CHAIN: B, D, F, H;	COMPLEX (ONCOGENE PROTEIN/PEPTIDE) C-CRK (N-TERMINAL SH3 DOMAIN) (C-CRKSH3-N) COMPLEXED WITH ICKA 3 C3G PEPTIDE (PRO-PRO-PRO-ALA-LEU- PRO-PRO-LYS-LYS-ARG)	PHOSPHOTRANSFERASE C- SRC KINASE (SH3 DOMAIN) (E.C.2.7.1.112) 1CSK 3	SYNTAXIN-1A; CHAIN: A, B, C;	SSOI PROTEIN; CHAIN: A;	PHOSPHOTRANSFERASE FYN: CHAIN: A; 3BP-2; CHAIN: B;
SeqFold score	66.14								
PMF		0.75	-	0.95	66.0	0.99	0.05	0.05	0.88
Verify		0.64	0.47	0.67	0.58	0.72	0.22	0.24	0.02
PSI- BLAST	0.0042	1.70E-08	4.20E-19	7.00E-18	8.40E-18	9.80E-19	2.80E-06	0.0056	2.80E-17
End	278	440	442	444	442	442	214	256	444
Start AA	88	370	388	390	389	387	93	166	385
Chain ID	4		<	<	<	<	V	A	٧
PDB ID	[avi	lavej	lazc	166z	lcka	lcsk	lez3	1fio	lfyn
SEQ TD	493	493	493	493	493	493	493	493	493

SEQ ID NO:	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
										TYROSINE-PROTEIN KINASE, 2 PHOSPHORYLATION, ATP-BINDING, MYRISTYLATION, SH3 DOMAIN, 3 COMPLEX (PHOSPHOTRANSFERASE/PEPTIDE)
493	1 g bq	¥	388	442	2.80E-18	0.34			GRB2; CHAIN: A; SOS-1; CHAIN: B;	COMPLEX (SIGNAL TRANSDUCTION/PEPTIDE) COMPLEX (SIGNAL TRANSDUCTION/PEPTIDE), SH3 DOMAIN
493	1gbr	₹	380	444	5.60E-19	0.47	_		SIGNAL TRANSDUCTION PROTEIN GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2, N-TERMINAL IGBR 3 SH3 DOMAIN) COMPLEXED WITH SOS-A PEPTIDE IGBR 4 (NMR, 29 STRUCTURES) IGBR 5	
493	1gri	V	272	444	1.10E-16	0.34	0.43		GROWTH FACTOR BOUND PROTEIN 2; IGRI 5 CHAIN: A, B; IGRI 6	SIGNAL TRANSDUCTION ADAPTOR SH2, SH3 IGRI 14
493	Thsq		383	444	2.80E-18	0.6	66:0		PHOSPHORIC DIESTER HYDROLASE PHOSPHOLIPASE C-GAMMA (SH3 DOMAIN) (E.C.3.1.4.11) 1HSQ 3 (NMR, MINIMIZED MEAN STRUCTURE) 1HSQ 4	
493	Iqly	⋖	389	444	4.20E-18	0.47	86.0		TYROSINE-PROTEIN KINASE BTK; CHAIN: A;	TYROSINE-PROTEIN KINASE BRUTONS TYROSINE KINASE, B CELL PROGENITOR KINASE, TRANSFERASE, TYROSINE-PROTEIN KINASE, PHOSPHORYLATION, 2 SH3 DOMAIN
493	1dde	⋖	-	262	4.20E-07			69.94	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	PROTEIN TRANSPORT HELIX-TURN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT
493	4hck		385	444	7.00E-19	0.1	0.99		HEMATOPOIETIC CELL KINASE; CHAIN: NULL;	TRANSFERASE HCK; SH3, PROTEIN TYROSINE KINASE, SIGNAL TRANSDUCTION, 2 TRANSFERASE
498	lath	4	342	422	5.60E-35			82.45	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC

PDB Chain	.E	Start	End	PSI- BLAST	Verify	PMF score	SeqFold	Coumpound	PDB annotation
	1							OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	FINGER, DNA-BINDING PROTEIN
Iallı A	1	370	478	5.60E-35	0.03			QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
lath A	1	398	\$05	1.40E-33	0.12	0.77		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
Ime V		257	338	6.80E-49	0.38	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGERIDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGERIDNA)
y De C		285	366	1.70E-50	0.4	-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION. PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
J C		313	394	3.40E-51	0.24			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN, CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Ine C	1	341	422	5.10E-51	0.34	_		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
C L me	ĺ	341	423	3.40E-51			110.98	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Jme C		397	477	3.40E-47	0.05	_		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Ime C		397	505	4.20E-36	-0.05	0.87		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC

	X, 2	INC X	INC X	NC X X	NOT	NOT	NOT	
uo	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX GYNC FINGEP DNAA	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION NITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POL YMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	rion MPLEX
PDB annotation	N-DNA PROTEIN CTURE, (NA)	CTURE, (NA)	PENGER IN-DNA PROTEIN CTURE, (PENGER N-DNA PROTEIN CTURE,	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRI INITIATION, ZINC FINGER PRO	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRI INITIATION, ZINC FINGER PRO	COMPLEX (TRANSCRIPTION REGUL ATION/DNA) COMPLEX (TRANSCRIPTION REGUL ATION/DNA), RNA POLYMERASE III, 2 TRANSCRI INTIATION, ZINC FINGER PRC	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX TRANSCRIPTION
PDB	FINGER, PROTEIN-DNA INTERACTION, PROTEII CRYSTAL STRUCTURE, (ZINC FINGER/DNA)	COMPLEX (ZINC FINGE FINGER, PROTEIN-DNA INTERACTION, PROTEII CRYSTAL STRUCTURE,	COMPLEX (ZING FINGE FINGER, PROTEIN-DNA FINGER, PROTEIN-DNA FINGER/CRYSTAL STRUCTURE, (ZING FINGER/DNA)	COMPLEX (ZINC FINGE FINGER, PROTEIN-DNA INTERACTION, PROTEII CRYSTAL STRUCTURE, (ZINC FINGER/DNA)	COMPLEX (TRANS REGULATION/DN. (TRANSCRIPTION REGULATION/DN. POLYMERASE III,	COMPLEX (TRAN) REGULATION/DN (TRANSCRIPTION REGULATION/DN REGULATION/DN POLYMERASE III,	COMPLEX (TRANK REGULATION/DN (TRANSCRIPTION REGULATION/DN POLYMERASE III,	COMPLEX (TRANS REGULATION/DN, (TRANSCRIPTION
	FINGER, INTERA CRYSTA (ZINC FI	COMPLE FINGER INTERA CRYSTA	COMPLI COMPLI FINGER INTERA CRYSTA	COMPLI FINGER INTERA CRYSTA (ZINC F.	COMPL REGUL, (TRANS REGUL, POLYM INITIAT	COMPL) REGUL, (TRANS REGUL, POLYM INITIAT	COMPL REGUL (TRANS REGUL POLYM	COMPL REGUL,
	GER G,	g; GER , G;	GER GER , G,	GER GER	S Æ.	S. Ë.	S Ę.	S. Æ.
Coumpound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F. G;	TFIIIA: CHAIN: A. D: 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;
Conm	ENSUS Z EIN, CHA	CHAIN:	CHAIN: ENSUS Z EIN; CH/	CHAIN: ENSUS Z EIN; CH/	TFIIIA: CHAIN: A. RIBOSOMAL RNA CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, RIBOSOMAL RNA CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, RIBOSOMAL RNA CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, RIBOSOMAL RNA
	CONSE	DNA; CONSI	DNA; CONSI	DNA; CONSI PROTI	TFIIIA RIBOS CHAIN	RIBOS CHAIN	RIBOS	RIBOS
SeqFold score						102.44		
PMF		-	_	-0.15	10.0-		0.94	0.92
Verify		0.23	0.19	0.1	90.0		0.11	-0.02
PSI- BLAST		1.00E-49	1.70E-35	3.40E-12	1.00E-33	1.40E-68	1.70E-36	8.50E-39
End		505	511	282	375	447	437	487
Start		424	452	255	221	285	286	342
Chain ID		U	O	D	< V	<	<	V.
PDB LD	۶	lmc y	lme y	i me	1116	1116	1116	1tf6
SEQ ID		498	498	498	498	498	498	498

PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
Coumpound		TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1, CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI: CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA: CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score					88.23		
PMF		0.95	0.05			-	_
Verify score		0.14	-0.02	0.25		0.23	0.36
PSI- BLAST		8.50E-35	1.00E-30	1.70E-34	2.80E-56	2.80E-56	8.50E-35
End		507	338	366	395	394	394
Start AA		370	228	265	285	290	293
Chain ID		<	O	O	U	U	ပ
PDB ID		1116	lubd	lubd	pqn1	lubd	lubd
SEQ ID NO:		498	498	498	498	498	498

pund PDB annotation		ADENO- COMPLEX (TRANSCRIPTION RUS P5 REGULATION/DNA) YING-YANG 1; MENT DNA; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX ATRANSCRIPTION REGII ATION/DNA)				ROTEIN GLII; COMPLEX (DNA-BINDING CHAIN: C, ZINC FINGER, COMPLEX (DNA-BINDING) BINDING PROTEIN/DNA)	ROTEIN GLII; COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	ROTEIN GLII; COMPLEX (DNA-BINDING; CHAIN: C, PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	ROTEIN GLII; COMPLEX (DNA-BINDING; CHAIN: C, PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)
SeqFold Coumpound score		YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	97.78 ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
PMF		_	0.92	0.28		_	-	0.86	0.41
Verify score		O	-0.07	-0.26		0.24	0.36	0.14	0.17
PSI- BLAST		2.80E-56	6.80E-36	8.50E-33	2.80E-70	2.80E-70	3.40E-33	1.30E-63	4.20E-61
End		422	505	337	423	422	423	479	507
Start AA		311	405	200	285	290	293	313	341
Chain ID	i 	O	U	∢	<	<	⋖	⋖	∢
PDB ID		lubd	lubd	2gli	2gli	. 2gli	2gli	2gli	2gli
SEQ ID		498	498	498	498	498	498	498	498

		7	T	1				
PDB annotation	PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE SKP2 F-BOX; SKP1; SKP1, SKP2, F-BOX, LRR, LEUCINE-RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION			LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
Coumpound	CHAIN: A; DNA; CHAIN: C, D;	SKP2; CHAIN: A. C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	CYCLIN A/CDK2- ASSOCIATED P19; CHAIN: A, C; CYCLIN A/CDK2- ASSOCIATED P45; CHAIN: B, D;	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;
SeqFold score								
PMF score		0.52	0.53	0.16	0.01	0.53	69:0	0.81
Verify score		-0.78	-0.81	0.08	-0.37	-0.07	0.02	-0.55
PSI- BLAST		0.0007	2.80E-06	1.70E-06	1.40E-13	8.40E-17	3.40E-16	5.60E-11
End		57	57	907	305	323	316	307
Start		28	28	998	255	259	261	262
Chain ID		A	¥	<				V
PDB ID		1fqv	Ifsl	1fbv	Ibor	1chc	1chc	Ifbv
SEQ ID		499	499	500	501	501	501	501

									
PDB annotation	LIGASE CBI, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	MEMBRANE PROTEIN LECTIN-LIKE, NEUROBIOLOGY, CELL-CELL ADHESION, CELL-CELL 2 RECOGNITION, ALTERNATIVE SPLICING, MEMBRANE PROTEIN	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	TRANSFERASE DINUCLEOTIDE- BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN
Coumpound	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	RAGI; CHAIN: NULL;	RAGI; CHAIN: NULL;	NEUREXIN-I BETA; CHAIN: A, B, C, D, B, F, G, H;	NICOTINATE MONONUCLEOTIDE:5,6- CHAIN: A;	NICOTINATE MONONUCLEOTIDE:5,6- CHAIN: A;	NICOTINÁTE MONONUCLEOTIDE:5,6- CHAIN: A;	ALKALINE PROTEASE; 1KAP 4 CHAIN: P; 1KAP 5 TETRAPEPTIDE (GLY SER
SeqFold score									
PMF score	_	0.22	0.94	0.45	-0.11	-0.2	-0.18	-0.2	-0.2
Verify score	-0.24	-0.17	-0.11	-0.27	0.21	0.3	0.51	0.56	96.0
PSI- BLAST	1.70E-09	7.00E-13	8.40E-18	1.70E-09	1.40E-17	1.10E-23	1.30E-20	1.10E-23	7.00E-14
End AA	310	319	333	342	224	1380	1072	1241	1439
Start	263	259	239	263	50	1042	753	116	1076
Chain ID	e e	Y			A	⋖	V	¥	d.
PDB ID	1fbv	1825	lrmd	Пще	lc4r	1d0s	sop1	140s	lkap
SEQ ID	201	501	201	501	502	502	502	502	502

PDB annotation		ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN IKAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN IKAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PROTEASE; IKAP 6 CALCIUM BINDING PROTEIN IKAP 19	OUTER MEMBRANE PROTEIN OSMOPORIN; OUTER MEMBRANE PROTEIN, NON-SPECIFIC PORIN, OSMOPORIN, 2 BETA-BARREL, TRANSMEMBRANE	OUTER MEMBRANE PROTEIN OSMOPORIN; OUTER MEMBRANE PROTEIN, NON-SPECIFIC PORIN, OSMOPORIN, 2 BETA-BARREL, TRANSMEMBRANE	OUTER MEMBRANE PROTEIN OSMOPORIN; OUTER MEMBRANE PROTEIN, NON-SPECIFIC PORIN, OSMOPORIN, 2 BETA-BARREL, TRANSMEMBRANE			METAL BINDING PROTEIN BETA SANDWICH, CALCIUM-BINDING
PD	1KAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PR. IKAP 6 CALCIUM BINDING P. IKAP 19	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PR. IKAP 6 CALCIUM BINDING P	ZINC METALLOPROTEASE P. AERUGINOSA ALKALINE PRO IKAP 6 CALCIUM BINDING P IKAP 19	OUTER MEMBRANE OSMOPORIN; OUTEI PROTEIN, NON-SPEC OSMOPORIN, 2 BETA	OUTER MEMBRANE OSMOPORIN; OUTEI PROTEIN, NON-SPEC OSMOPORIN, 2 BETA	OUTER MEMBRANE OSMOPORIN; OUTE PROTEIN, NON-SPE OSMOPORIN, 2 BET TRANSMEMBRANE			METAL BINDIN SANDWICH, CA
Coumpound	ASN SER); 1KAP 9 CHAIN: 1; 1KAP 10	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER ASN SÉR); IKAP 9 CHAIN: I; IKAP 10	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER ASN SER); IKAP 9 CHAIN: I; IKAP 10	ALKALINE PROTEASE; IKAP 4 CHAIN: P; IKAP 5 TETRAPEPTIDE (GLY SER ASN SER); IKAP 9 CHAIN: 1; IKAP 10	OMPK36; CHAIN: A, B, C;	OMPK36; CHAIN: A, B, C;	OMPK36; CHAIN: A, B, C;	OUTER MEMBRANE PROTEIN PHOSPHOPORIN (PHOE) IPHO 3	OUTER MEMBRANE PROTEIN PHOSPHOPORIN (PHOE) 1PHO 3	CHAIN ALPHA2 CHAIN; CHAIN: A, B, C, D;
SeqFold score										
PMF score		-0.19	-0.2	-0.19	-0.2	-0.19	-0.2	-0.2	-0.19	0.1
Verify		=	1.03	0.87	1.07	1.04	0.87	0.83	1.15	0.51
PSI- BLAST		5.60E-10	8.40E-14	1.10E-13	1.10E-31	1.40E-32	4.20E-27	1.40E-23	7.00E-27	2.80E-13
End		743	858	1244	1348	1043	1241	1045	1214	225
Start AA		482	530	890	1004	707	875	692	872	78
Chain ID		a.	۵.	a.	⋖	<	<			⋖
PDB ID		lkap	1kap	lkap	losm	losm	losm	1 pho	1pho	0nb1
SEQ B NO:		502	502	502	502	502	502	502	502	502

PDB annotation	PROTEIN INTEGRAL MEMBRANE PROTEIN PORIN MATRIX PORIN, OMPF PORIN; 20MF 7 PORIN, MEMBRANE PROTEIN 20MF 12		TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN SIGNAL TRANSDUCTION PROTEIN	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN SIGNAL TRANSDUCTION PROTEIN CYTOSKELETON	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN SIGNAL TRANSDUCTION PROTEIN CYTOSKELETON	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN SIGNAL TRANSDUCTION PROTEIN CYTOSKELETON CYTOSKELETON CYTOSKELETON	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN SIGNAL TRANSDUCTION PROTEIN CYTOSKELETON CYTOSKELETON CYTOSKELETON CYTOSKELETON TYTOSKELETON CYTOSKELETON T	IRANSFERASE PROTEIN-ACETYL SOENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN SIGNAL TRANSDUCTION PROTEIN CYTOSKELETON CYTOSKELETON SYTOSKELETON SYTOSKELETON SYTOSKELETON SYTOSKELETON SYTOSKELETON SYTOSKELETON THE SYTOSKELETON SYTOSKELETON SYTOSKELETON SYTOSKELETON TRANSDUCTION PROTEIN, 3- PHOSPHOINOSITIDES, INOSITOL TRANSDUCTION PROTEIN, ADAPTOR PROTEIN PROTEIN	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE SIGNAL TRANSDUCTION PROTEIN CYTOSKELETON CYTOSKELETON CYTOSKELETON CYTOSKELETON CYTOSKELETON CYTOSKELETON TERANSDUCTION PROTEIN SIGNALING PROTEIN Jahry BAM32; PLECKSTRIN, 3- PHOSPHOINOSITIDES, INOSITOL TERANSDUCTION PROTEIN, ADAPTOR PROTEIN SIGNAL TRANSDUCTION SON OF SEVENLESS; PLECKSTRIN, SON OF SEVENLESS; PLECKSTRIN, SON OF SEVENLESS, SIGNAL TRANSDUCTION
		TRANSFER			. 4 4						
	MATRIX PORIN OUTER MEMBRANE PROTEIN F; 20MF 5 CHAIN: NULL; 20MF 6	HPA2 HISTONE	ACETYLTRANSFERASE; CHAIN: A, B, C, D;	ACETYLTRANSFERASE CHAIN: A, B, C, D; BETA-SPECTRIN; 1BTN CHAIN: NULL; 1BTN 5	ACETYLTRANSFERASE CHAIN: A, B, C, D; BETA-SPECTRIN; IBTN CHAIN: NULL; IBTN 5 BETA-SPECTRIN; IBTN CHAIN: NULL; IBTN 5	ACETYLTRANSFERASE; CHAM: A, B, C, D; BETA-SPECTRIN; IBTN 4 ECHAM: NULL; IBTN 5 ECHAM: NULL; IBTN 4 ECHAM: NULL; IBTN 5 BETA-SPECTRIN; IDRO 6 CHAM: NULL; IDRO 6	ACETYLTRANSFERASE; CHAIN: A, B, C, D; BETA-SPECTRIN; IBTN 5 CHAIN: NULL; IBTN 5 BETA-SPECTRIN; IBTN 5 BETA-SPECTRIN; IDRO CHAIN: NULL; IDRO 7 BETA-SPECTRIN; IDRO 7 CHAIN: NULL; IDRO 7	ACETYLTRANSFERASE CHAIN: A, B, C, D; BETA-SPECTRIN; IBTN 5 CHAIN: NULL; IBTN 5 BETA-SPECTRIN; IBTN 5 BETA-SPECTRIN; IDRO 7 BETA-SPECTRIN; IDRO 7 BETA-SPECTRIN; IDRO 7 BETA-SPECTRIN; IDRO 7 BETA-SPECTRIN; IDRO 7 BETA-SPECTRIN; IDRO 7 BETA-SPECTRIN; IDRO CHAIN: NULL; IDRO 7	ACETYLTRANSFERASE; CHAIN: A, B, C, D; BETA-SPECTRIN; 1BTN 4 CHAIN: NULL; 1BTN 5 BETA-SPECTRIN; 1BTN 6 CHAIN: NULL; 1BTN 6 CHAIN: NULL; 1DRO 7 BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7 BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7 BETA-SPECTRIN; 1DRO 6 CHAIN: NULL; 1DRO 7 DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A;	ACETYLTRANSFERASE; CHAIN: A, B, C, D; BETA-SPECTRIN; IBTN 4 CHAIN: NULL; IBTN 5 BETA-SPECTRIN; IBTN 4 CHAIN: NULL; IBTN 6 CHAIN: NULL; IBTN 6 CHAIN: NULL; IDRO 7 BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7 BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7 BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7 BETA-SPECTRIN; IDRO 6 CHAIN: NULL; IDRO 7 DUAL ADAPTOR OF PHOSPHOTYROSINE AND 3- CHAIN: A; PHOSPHORYLATION PLECKSTRIN (N-TERMINAL PLECKSTRIN (N-TERM	ACETYLTRANSFERAS CHAIN: A, B, C, D; BETA-SPECTRIN, 1BTN CHAIN: NULL; 1BTN S BETA-SPECTRIN, 1BTN CHAIN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THAN: NULL; 1DRO THOSPHORYLATION PLECKSTRIN (A-TER) PLECKSTRIN (A-TER) PLECKSTRIN (A-TER) PLECKSTRIN (A-TER) THE U GLU (HIS) ADDED TO THE C TERMINUS IPLS 4 (IN) LEHHHHHHH) (NMR, 2 STRUCTURES) 1PLS 5 SOS 1; CHAIN: NULL;
	MA MEI 20N 6	Tal.	ACE CH/	ACE CH/ CH/ CH/ CH/	A CCHA						
e score						\$0.76	50.76	50.76			
score score	-0.18	90.0		0.86	0.86	0.86	0.86	0.78	0.39 0.75 0.75 0.75	0.78 0.39 0.75 -0.07	0.78 0.39 0.39 0.16 0.16
score		-0.12		0.36	0.36	0.36	0.36	0.36 0.38 0.08	0.36 0.08 0.35 0.15	0.36 0.08 0.15 0.05	0.36 0.38 0.35 0.15 0.05
BLAST	7.00E-21	0.0028		1.70E-22	1.70E-22 2.80E-24	1.70E-22 2.80E-24 7.00E-28	1.70E-22 2.80E-24 7.00E-28 5.10E-17	1.70E-22 2.80E-24 7.00E-28 5.10E-17 7.00E-28	1.70E-22 2.80E-24 7.00E-28 5.10E-17 7.00E-28 3.40E-17	1.70E-22 2.80E-24 7.00E-28 5.10E-17 7.00E-28 3.40E-17	1.70E-22 2.80E-24 7.00E-28 5.10E-17 7.00E-28 3.40E-17 1.70E-18
AA	1202	282		187	187	187	187 187 192 190	187 187 192 190 189	187 192 190 189 185	187 192 198 189 188 188	187 192 198 189 189 190
VΥ	842	227		82	82	82 83 73	82 83 73	82 83 73 83	82 83 73 83 88	88 88 88 88	88 88 88 88 88 88
a		4									
3	2omf	Idsm	-	1 btn	1btn 1btn	1btn 1btn 1dro	lbtn lbtn ldro	lbtn ldro ldro	 		
e ë	502	206		507	507	507	507 507 507 507	507 507 507 507	\$07 \$07 \$07 \$07 \$07	\$07 \$07 \$07 \$07 \$07 \$07	\$07 \$07 \$07 \$07 \$07 \$07 \$07 \$07

	z	40LOGY	ONIN IN 2		IA ILS,	NSEC1;		NSEC1;	NOED I			ED 35	IELIX	LE- CTILE		ONE,	JNG,	ONE, ING,
PDB annotation	BINDING, PHOSPHORYLATION	STRUCTURAL PROTEIN DYSTROPHIN, MUSCULAR DYSTROPHY, CALPONIN HOMOLOGY DOMAIN, 2 ACTIN-BINDING, UTROPHIN	STRUCTURAL PROTEIN CALPONIN HOMOLOGY DOMAIN, DOMAIN SWAPPING, ACTIN BINDING, 2 UTROPHIN, DYSTROPHIN, STRUCTURAL PROTEIN	OWE INTERNATION	SIRUCLIURAL FRUIEIN I WO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS,	ENDOCYTOSIS/EXOCYTOSIS NSECI;	PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX, MIII TI-SURI NIT	FNDOCYTOSIS/FYDCYTOSIS NGFCI	ENDOCT TOSISTENCE TOSIST PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS	SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BINDI F	MEMBRANE PROTEIN FOUR HELIX BUNDLE, ALPHA HELIX	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE	NI	CHAPERONE HSP40; CHAPERONE,	HEAT SHOCK, PROTEIN FOLDING, DNAK	CHAPERONE HSP40, CHAPERONE, HEAT SHOCK, PROTEIN FOLDING,
	BINDI	STRUCTUR DYSTROPH DYSTROPH DOMAIN, 2 UTROPHIN	STRUC HOMC SWAP UTRO STRUC	7110110	REPE/ HELIC TAND	ENDO	MULT	ENDO PROTI	FNDO	PROTI	ENDO	SYNAPTC KDA PRO	MEME	CONT	PROTEIN	CHAP	HEAT	CHAPI HEAT
Coumpound		DYSTROPHIN; CHAIN: A, B, C, D;	UTROPHIN ACTIN BINDING REGION; CHAIN: A, B;	AT DITA COPOCEDIO ATTAIN	ALFRA SFECTIVIII, CHAIIN: A, B, C;	SYNTAXIN BINDING	PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A: CHAIN: B:	SYNTAXIN BINDING	PROTEIN I; CHAIN: A;	SYNTAXIN-1A; CHAIN: A, B,	ပ်	SSOI PROTEIN; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN:	A;	DNAJ; CHAIN: NULL;		DNAJ; CHAIN: NULL;
SeqFold score					<u>.</u>											57.77		
PMF		0.03	0.27	000	60.0	0.05		0.3	0.03		0.03		0.18	90.0			•	1
Verify score		-0.01	0.01	70.0	9	-0.13		-0.31	-0.3	C	0.25		-0.45	-0.12				0.45
PSI- BLAST		8.50E-30	1.70E-29	00 300 0	4.00E-03	1.40E-18		1.10E-10	1.10F-08		1.10E-09		2.80E-05	4.20E-21		6.80E-33		6.80E-33
End		248	248	673	700	460		561	940		933		686	467		124		122
Start		106	011	376		208		364	713	}	804		794	205		46		47
Chain ID		<	4	_	¢	В		æ	В	1	∢		٧	V				
PDB ID		Ідхх	lqag	-		1db1		1dn1	Idn1		lez3		l fio	Iquu		1bq0		1bq0
SEQ ID NO:		508	508	71.5		515		515	515		515		515	515		516		516

	-т			 																		Γ	\neg	Τ-			T		
PDB annotation	DNAK	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE									SUGAR BINDING PROTEIN BETA	TREFOIL, MULTILECTIN RECEPTOR,	FILULIARY HORMONES, 2 SULFALED CARBOHYDRATE	TRANSFERASE	GLYCOSYLTRANSFERASE		TRANSFERASE	GLYCOSYLTRANSFERASE		HYDROLASE XYLAN DEGRADATION								
Coumpound		HUMAN HSP40; CHAIN: NULL;	HUMAN HSP40; CHAIN: NULL;	COMPLEX	RATE) ABRIN-A	COMPLEXED WITH TWO	SUGAR CHAINS 1ABR 3	COMPLEX	(GLYCOSIDASE/CARBOHYD	KATE) ABRIN-A COMPLEXED WITH TWO	SUGAR CHAINS LABR 3	MANNOSE RECEPTOR;	CHAIN: A;		SPORE COAT	POLYSACCHARIDE	BIOSYNTHESIS PROTEIN CHAIN: A;	SPORE COAT	POLYSACCHARIDE	CHAIN: A;	ENDO-1,4-BETA-XYLANASE; CHAIN: A. B:	GLYCOSIDASE RICIN	(E.C.3.2.2.22) 2AAI 3	OX VGEN TRANSPORT	HEMOGLOBIN (DEOXY,	HUMAN FETAL F=/IIS=)	OXYGEN TRANSPORT	HEMOGLOBIN (DEOXY,	HUMAN FETAL F=/(II \$=) 1FDHG 1 1FDHH 2
SeqFold		52.89																						117.26	27:3:				
PMF			-	0.01				0.04				0.13			0.11			0	_		96:0	0.03					-	•	
Verify			0.21	-0.08				0.04				-0.22			0.17			-0.1			0.08	0.07					0.70	7.0	
PSI- BLAST		1.00E-30	1.00E-30	1.40E-10				6.80E-30				0.0017			5.10E-23			7.00E-45			1.70E-31	1.70E-28		1 005,30	200:		1 00E-30	(C-700:1	
End		125	122	547				547				513			333			371			548	547		8	:		6	<u> </u>	
Start		46	48	378				412				444			Ξ			Ξ			422	413			•		-		
Chain ID				æ				В				A			A			V			V	В		c	·		ŀ	,	
PDB ID		1hdj	1hdj	labr		-		labr				1dqg	!		lqga			lqgq			lxyf	2aai		1 Cdh			1 fdh		
SEQ ID		516	516	522				522				522			522			522			522	522		531	3		\$73	77	

	PDB ID	Chain	Start AA	End AA	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
	Igcv	В	-	92	7.00E-40	90:0	-		HEMOGLOBIN; CHAIN: A, C; HEMOGLOBIN; CHAIN: B, D;	OXYGEN STORAGE/TRANSPORT HEMOGLOBIN, DEOXY FORM
	lawj		1018	1088	1.40E-18	0.45	0.15		ITK; CHAIN: NULL;	TRANSFERASE IL-2-INDUCIBLE T- CELL KINASE; TRANSFERASE, REGULATORY INTRAMOLECULAR
·+	laze	A	1037	1087	2.80E-18	0.5	0.89		GRB2; CHAIN: A; SOS; CHAIN: B:	COMPLEX, AINAGE COMPLEX (ADAPTOR PROTEIN/PEPTIDE) ASH, GROWTH 5: COTOR RECEPTOR-BOUND PROTEIN 7: COMPLEX (ADAPTOR PROTEIN/PEPTIDE), SH3 DOMAIN, 2 FACTOR FACTOR
+	1671	4	-	739	0			335.35	MYOSIN HEAVY CHAIN; CHAIN: A; MYOSIN REGULATORY LIGHT CHAIN; CHAIN: Y; MYOSIN ESSENTIAL LIGHT CHAIN;	MYOSIN MYOSIN MOTOR
 	1b7t	A	-	739	0	0.34	-		MYOSIN HEAVY CHAIN; CHAIN: A; MYOSIN REGULATORY LIGHT CHAIN; CHAIN: Y; MYOSIN ESSENTIAL LIGHT CHAIN;	MYOSIN MYOSIN MOTOR
+	1br1	4	-	708	0	0.59	_		MYOSIN, CHAIN: A, B, C, D, E, F, G, H;	MUSCLE PROTEIN MDE; MUSCLE PROTEIN
	Ibri	¥.	_	711	0			352.11	MYOSIN; CHAIN: A, B, C, D, E, F, G. H;	MUSCLE PROTEIN MDE, MUSCLE PROTEIN
т -	1br2	⋖	_	629	0	ļ 		339.22	MYOSIN; CHAIN: A, B, C, D, E, F;	MUSCLE PROTEIN MUSCLE PROTEIN
$\overline{}$	1672	<		629	0	0.44	-		MYOSIN; CHAIN: A, B, C, D, E, F;	MUSCLE PROTEIN MUSCLE PROTEIN
	善	¥	_	739	0	0.01	_		MYOSIN HEAD, CHAIN: A; MYOSIN HEAD, CHAIN: Y; MYOSIN HEAD, CHAIN: Z;	CONTRACTILE PROTEIN MYOSIN MOTOR, CONFORMATIONAL CHANGES
Γ	1gbr	¥	1028	1089	1.10E-19	0.45	0.72		SIGNAL TRANSDUCTION PROTEIN GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2, N-TERMINAL IGBR	

							· · · · · · · · · · · · · · · · · · ·	
PDB annotation			SIGNAL TRANSDUCTION ADAPTOR SH2, SH3 IGRI 14		CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, A TP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN- BINDING, ATP-BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN
Coumpound	3 SH3 DOMAIN) COMPLEXED WITH SOS-A PEPTIDE IGBR 4 (NMR, 29 STRUCTURES) IGBR 5	ADAPTOR PROTEIN CONTAINING SH2 AND SH3 GROWTH FACTOR RECEPTOR-BOUND PROTEIN 2 (GRB2) IGFC 3 (C- TERMINAL SH3 DOMAIN) (NMR, MINIMIZED MEAN STRUCTURE) IGFC 4	GROWTH FACTOR BOUND PROTEIN 2; 1GRI 5 CHAIN: A, B; 1GRI 6	PHOSPHORIC DIESTER HYDROLASE PHOSPHOLIPASE C-GAMMA (SH3 DOMAIN) (E.C.3.1.4.11) 11SQ 3 (NMR, MINIMIZED MEAN STRUCTURE) 11SQ 4	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;
SeqFold score						348.12	269.23	
PMF score		0.59	0.12	1202.08	-			
Verify score		0.29	-0.23	0.24	0.18			0.37
PSI- BLAST		1.30E-19	8.40E-17	1.40E-18	0	0	0	0
End		1089	1089	1089	678	629	809	809
Start AA		1035	896	1032	_	-	-	_
Chain ID			¥					
PDB ID		lgfc	lgri	lhsq	IIvk	livk	lmn d	lmn d
SEQ NO:		526	526	526	526	526	526	526

PDB ID		Chain	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
lpwt	1		1032	1089	7.00E-20	29.0	-1.41		ALPHA SPECTRIN; CHAIN: NULL;	CIRCULAR PERMUTANT PWT; CIRCULAR PERMUTANT, SH3 DOMAIN, CYTOSKELETON
1qly		<	1036	1089	4.20E-18	0.25	-1.41		TYROSINE-PROTEIN KINASE BTK; CHAIN: A;	TYROSINE-PROTEIN KINASE BRUTONS TYROSINE KINASE, B CELL PROGENITOR KINASE, TRANSFERASE, TYROSINE-PROTEIN KINASE, PHOSPHORYLATION, 2 SH3 DOMAIN
Isem	E	∢	1035	6801	4.20E-19	0.63	0.94		SEM-5; 1SEM 3 CHAIN: A, B; 1SEM 5 10-RESIDUE PROLINE-RICH PEPTIDE FROM MSOS 1SEM 8 CHAIN: C, D 1SEM 10	SIGNAL TRANSDUCTION PROTEIN SRC-HOMOLOGY 3 (SH3) DOMAIN, PEPTIDE-BINDING PROTEIN, ISEM 18 2 GUANINE NUCLEOTIDE EXCHANGE FACTOR ISEM 19
2mys	sx	<	_	731	0	-0.02	-		MYOSIN; CHAIN: A, B, C;	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN
EZ.	2mys	Y	1	737	0			267.23	MYOSIN; CHAIN: A, B, C;	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN
4hck	×		1032	1089	1.10E-18	0.32	0.18		HEMATOPOIETIC CELL KINASE; CHAIN: NULL;	TRANSFERASE HCK; SH3, PROTEIN TYROSINE KINASE, SIGNAL TRANSDUCTION, 2 TRANSFERASE
3grx	ř		14	96	0.0015	-0.21	0.68		GLUTAREDOXIN 3; CHAIN: NULL;	ELECTRON TRANSPORT ELECTRON TRANSPORT, THIOL-DISULFIDE OXIDOREDUCTASE, 2 THIOLTRANSFERASE, THIOREDOXIN SUPERFAMILY
3grx	×		23	68	9.80E-06	0.17	0.96		GLUTAREDOXIN 3; CHAIN: NULL;	ELECTRON TRANSPORT ELECTRON TRANSPORT, THIOL-DISULFIDE OXIDOREDUCTASE, 2 THIOLTRANSFERASE, THIOREDOXIN SUPERFAMILY
1=	1991	8	14	159	3.40E-19	0.25	96:0		ARYLALKYLAMINE N- ACETYLTRANSFERASE; CHAIN: A, B;	TRANSFERASE ACETYLTRANSFERASE
2	1587	¥	24	184	6.80E-08	0.29	0.95		AMINOGLYCOSIDE N6'- ACETYLTRANSFERASE TYPE 1; CHAIN: A;	TRANSFERASE AAC; AMINOGLYCOSIDE 6:-N- ACETYLTRANSFERASE, ANTIBIOTIC

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold	Coumpound	PDB annotation
										2 RESISTANCE, ACETYL COENZYME A
529	lcjw	4	14	159	1.20E-19	0.5	0.88		SEROTONIN N- ACETYLTRANSFERASE; CHAIN: A;	TRANSFERASE N-ACETYL TRANSFERASE
529	lcm 0	В	36	174	2.80E-13	0.39	0.81		P300/CBP ASSOCIATING FACTOR; CHAIN: B, A;	SIGNALING PROTEIN P300/CBP ASSOCIATED FACTOR, COENZYME A, ACETYLTRANSFERASE, 2 COACTIVATOR, SIGNALING PROTEIN
529	lcm 0	В	7.1	184	1.70E-05	0.15	0.19		P300/CBP ASSOCIATING FACTOR; CHAIN: B, A;	SIGNALING PROTEIN P300/CBP ASSOCIATED FACTOR, COENZYME A, ACETYLTRANSFERASE, 2 COACTIVATOR, SIGNALING PROTEIN
529	Iqsm	<	=	156	1.50E-13	0.13	0.33		HPA2 HISTONE ACETYLTRANSFERASE; CHAIN: A, B, C, D;	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE
529	lqsm	<	12	165	4.20E-19	0.15	0.23		HPA2 HISTONE ACETYLTRANSFERASE; CHAIN: A, B, C, D;	TRANSFERASE PROTEIN-ACETYL COENZYME A COMPLEX, ACETYLTRANSFERASE
529	lqst	<	79	183	3.40E-07	90:0-	61.0		TGCNS HISTONE ACETYL TRANSFERASE; CHAIN: A;	TRANSFERASE HISTONE ACETYLTRANSFERASE, GCN3- RELATED N-ACETYLTRANSFERASE, 2 COA BINDING PROTEIN
529	lygh	∢	80	179	1.00E-05	0.09	0.04		TRANSCRIPTIONAL ACTIVATOR GCN5; CHAIN: A, B;	GENE REGULATION ADA4; TRANSCRIPTIONAL REGULATION, HISTONE ACETYLATION, N-2 ACETYLTRANSFERASE, GCN5 RELATED N-ACETYLTRANSFERASE FAMILY, 3 GENE REGULATION
534	lc1g	<	£	296	1.50E-54	-0.48	-		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
534	lclg	<	E	245	3.40E-49	-0.17	0.64		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
534	1c1g	4	3	248	5.10E-53	-0.17	0.94		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
535	lclg	4	3	296	1.50E-54	-0.48			TROPOMYOSIN; CHAIN: A,	CONTRACTILE PROTEIN

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
NO:	a l	a	۸A	ΑA	BLASI	score	score	score		
									B, C, D	TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
535	lclg	<	3	245	3.40E-49	-0.17	0.64		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
535	lclg	4	3	248	5.10E-53	-0.17	0.94		TROPOMYOSIN; CHAIN: A, B, C, D	CONTRACTILE PROTEIN TROPOMYOSIN COILED-COIL ALPHA- HELICAL, CONTRACTILE PROTEIN
538	la4y	A	9	306	4.20E-25	0.29	0.47		RIBONUCLEASE INHIBITOR, CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS
538	la9n	¥	136	297	2.80E-22	0.75	-		U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
538	1a9n	٧	164	306	4.20E-18	0.42	0.45		U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D:	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
538	la9n	«	210	284	0.00017	0.59	0.29		U2 RNA HAIRPIN IV; CHAIN; Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D:	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
238	1a9n	4	40	144	3.40E-06	-0.09	0.12		U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
538	la9n	ပ	136	301	5.60E-23	0.65	6.0		U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
538	Та 9п	ပ	210	284	0.00017	0.48	0.29		U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B": CHAIN: B, D:	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
538	1a9n	O	40	144	3.40E-06	0.1	0.03		U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTEIN
538	90P1	4	127	286	1.70E-22	0.41	8.0		INTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION
538	1d0b	Y	183	350	1.00E-16	0.19	0.21		INTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE FUCH REPEAT, CALCIUM BINDING, CELL

Chain Start E ID AA A		E V	End	PSI- BLAST	Verify score	PMF score	SeqFold	Coumpound	PDB annotation
3 162 5.10E-24	162		5.10E-2	4	0.3	0.11		INTERNALIN B; CHAIN: A;	ADHESION CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION
32 191 1.00E-25	161	 	1.00E-25		0.05	0.15		INTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION
57 212 8.50E-25	212		8.50E-25		0.17	0.22		NTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION
7 130 1.20E-09	130	 	1.20E-09		0.02	-0.14		RAB GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT; CHAIN: A, C; RAB GERANYLGERANYLTRANSF ERASE BETA SUBUNIT; CHAIN: B, D.	TRANSFERASE CRYSTAL STRUCTURE, RAB GERANYLGERANYLTRANSFERASE, 2.0 A 2 RESOLUTION, N- FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT
16 143 6.80E-13	143		6.80E-13		-0.03	0.16		OUTER ARM DYNEIN: CHAIN: A:	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER. DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA
52 120 1.00E-06	120		1.00E-06	1	-0.32	0.13		NUCLEAR RNA EXPORT FACTOR 1; CHAIN: A, B;	RNA BINDING PROTEIN TAP (NFX1), RIBONUCLEOPROTEIN (RNP,RBD OR RRM) AND LEUCINE-RICH-REPEAT 2 (LRR)
227 290 1.70E-07	290		1.70E-07		-0.1	0.03		NUCLEAR RNA EXPORT FACTOR 1; CHAIN: A, B;	RNA BINDING PROTEIN TAP (NFX1), RIBONUCLEOPROTEIN (RNP,RBD OR RRM) AND LEUCINE-RICH-REPEAT 2 (LRR)
52 120 1.00E-06	120	-	1.00E-06		-0.41	0.1		NUCLEAR RNA EXPORT FACTOR 1; CHAIN: A, B;	RNA BINDING PROTEIN TAP (NFX1); RIBONUCLEOPROTEIN (RNP,RBD OR RRM) AND LEUCINE-RICH-REPEAT 2 (LRR)
111 284 5.10E-13	284		5.10E-13		0.63	0.49		SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45, CYCLIN A/CDK2-ASSOCIATED PROTEIN P19, SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE
9 339 2.80E-17	339		2.80E-17		80.0	-0.09		SKP2; CHAIN: A, C, E, G, I, K,	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45: CYCLIN

SEQ	PDB	Chain ID	Start	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
Ö									H, J, L, N, P;	A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT. SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE
538	162	4	=	284	5,10E-13	0.13	0.24		SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE
538	1fs2	Y	84	306	1.30E-23	0.29	0		SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE
538	lyrg	∢	98	291	2.80E-22	0.26	-0.11		GTPASE-ACTIVATING PROTEIN RNA1_SCHPO; CHAIN: A, B;	TRANSCRIPTION RNA IP; RANGAP; GTPASE-ACTIVATING PROTEIN FOR SPII, GTPASE-ACTIVATING PROTEIN, GAP, RNA IP, RANGAP, LRR, LEUCINE-2 RICH REPEAT PROTEIN, TWINNING, HEMIHEDRAL TWINNING, MEROHEDRAL TWINNING, MEROHEDRAY
538	2bnh		63	311	1.40E-32	0.43	0.22		RIBONUCLEASE INHIBITOR; CHAIN: NULL;	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS
539	1d2г	<	35	359	0	0.54	_		TRYPTOPHANYL TRNA SYNTHETASE: CHAIN: A, B, C, D, E, F;	LIGASE TRPRS; CLASS I TRNA SYNTHETASE, AARS, INDUCED FIT, TRPRS
540	1406	4	274	371	4.20E-17	0.23	0.23		NITROGEN FIXATION REGULATORY PROTEIN FIXL; CHAIN: A;	SIGNALING PROTEIN OXYGEN SENSOR, HISTIDINE KINASE, PAS, HIGH-RESOLUTION, TWO- 2 COMPONENT SYSTEM
541	laew		30	199	7.00E-84			194.39	FERRITIN; CHAIN: NULL;	IRON STORAGE IRON STORAGE, MULTIGENE FAMILY, ACETYLATION
541	lacw		30	199	7.00E-84	0.36	_		FERRITIN; CHAIN: NULL;	IRON STORAGE IRON STORAGE,

PDB annotation	MULTIGENE FAMILY, ACETYLATION	IRON STORAGE IRON STORAGE, DIIRON	IRON STORAGE IRON STORAGE. DIIRON	IRON STORAGE IRON STORAGE	IRON STORAGE IRON STORAGE	IRON STORAGE IRON STORAGE	ISOMERASE ISOMERASE, PPIASE	COMPLEX (ISOMERASEPEPTIDE) COMPLEX (ISOMERASEPEPTIDE), CYCLOPHILIN A, HIV-1 CAPSID, 2 PSEUDO-SYMMETRY	COMPLEX (ISOMERASE/PEPTIDE) COMPLEX (ISOMERASE/PEPTIDE), CYCLOPHILIN A, HIV-1 CAPSID, 2 PSEUDO-SYMMETRY			COMPLEX (ISOMERASE/IMMUNOSUPPRESSANT) CYCLOSPORIN, ISOMERASE, ROTAMASE, SIGNAL ICYN 19	COMPLEX (ISOMERASE/IMMUNOSUPPRESSANT) CYCLOSPORIN, ISOMERASE, ROTAMASE, SIGNAL 1CYN 19	ISOMERASE(PEPTIDYL-PROLYL CIS- TRANS) PEPTIDYL-PROLYL CIS-
Coumpound		M FERRITIN; CHAIN: A.B.C.D.E.F.G.H.I.J.K.L.M.N.O P.Q.R.S.T.U.V.W.X;	M FERRITIN: CHAIN: A,B,C,D,E,F,G,H,I,J,K,L,M,N,O P,Q,R,S,T,U,V,W,X;	FERRITIN; CHAIN: NULL;	FERRITIN; CHAIN: NULL;	FERRITIN; CHAIN: NULL;	CYCLOPHILIN; CHAIN: NULL;	CYCLOPHILIN A; CHAIN: A; PEPTIDE FROM THE HIV-1 CAPSID PROTEIN; CHAIN: B;	CYCLOPHILIN A; CHAIN: A; PEPTIDE FROM THE HIV-1 CAPSID PROTEIN; CHAIN: B;	ISOMERASE(PEPTIDYL- PROLYL CIS-TRANS) CYCLOPHILIN (NMR, 12 STRUCTURES) ICLH 3	ISOMERASE(PEPTIDYL- PROLYL CIS-TRANS) CYCLOPHILIN (NMR, 12 STRUCTURES) ICLH 3	CYCLOPHILIN B: ICYN 6 CHAIN: A; ICYN 7 [D- (CHOLINYL)ALA]8- CYCLOSPORIN; ICYN 10 CHAIN: C; ICYN 11	CYCLOPHILIN B, ICYN 6 CHAIN: A, ICYN 7 [D- (CHOLINYL)ALA]8- CYCLOSPORIN; ICYN 10 CHAIN: C, ICYN 11	CYCLOPHILIN 3; CHAIN: A;
SeqFold score		221.42		284.09			94.49	89.5		9.69		103.11		
PMF			_			-			-		0.46		-	-
Verify score			0.53		0.5	0.5			0.7		0.28		0.52	0.62
PSI- BLAST		5.60E-88	5.60E-88	1.40E-75	1.40E-75	1.00E-74	5.10E-29	5.10E-38	5.10E-38	8.40E-39	8.40E-39	5.10E-34	5.10E-34	1.70E-34
End		199	661	200	200	200	162	163	160	991	163	171	160	160
Start		29	30	29	29	29	_	-	2	_	E .	_	2	2
Chain ID		¥	<					V	<			Y	<	∢
PDB ID		lmfr	lmfr	2fha	2fha	2fha	1a58	lawq	lawq	-1clh	1clh	lcyn	lcyn	1dy w
SEQ 1D		541	541	541	541	541	546	546	546	546	546	546	546	546

-	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
										TRANS ISOMERASE 3, ISOMERASE, ROTAMASE
=	1lop	V.	2	163	7.00E-41			79.69	CYCLOPHILIN A; CHAIN: A; SUCCINYL-ALA-PRO-ALA-P- NITROANILIDE; CHAIN: B;	COMPLEX (ISOMERASE/PEPTIDE) ISOMERASE, ROTAMASE, COMPLEX (ISOMERASE/PEPTIDE)
=	11op	A	3	162	7.00E-41	0.47	86.0		CYCLOPHILIN A; CHAIN: A; SUCCINYL-ALA-PRO-ALA-P- NITROANILIDE; CHAIN: B;	COMPLEX (ISOMERASE/PEPTIDE) ISOMERASE, ROTAMASE, COMPLEX (ISOMERASE/PEPTIDE)
اے	Iqng	<	2	160	6.80E-33	0.56	-		CYCLOPHILIN; CHAIN: A; CYCLOSPORIN A; CHAIN: D;	PEPTIDYLPROLYL CIS-TRANS ISOMERASE; CYCLOPHILIN A, CYCLOSPORIN A, PEPTIDYL CIS- TRANS ISOMERASE
	1qoi	4	2	160	1.70E-31	0.57	-		SNUCYP-20, CHAIN: A;	ISOMERASE USA-CYP; SNUCYP-20, CYCLOPHILIN, SNRNP, SPLICEOSOMAL
<u> </u>	1qoi	<	7	156	2.80E-36	0.31	0.98		SNUCYP-20; CHAIN: A;	ISOMERASE USA-CYP, SNUCYP-20, CYCLOPHILIN, SNRNP, SPLICEOSOMAL
100	2ттс	<	2	160	3.40E-31	9.0	-		COMPLEX (ISOMERASE/IMMUNOSUPP RESSANT) CYCLOPHILIN C COMPLEXED WITH CYCLOSPORIN A 2RMC 3	
[7]	2ттс	<	2	172	8.40E-41			94.18	COMPLEX (ISOMERASE/IMMUNOSUPP RESSANT) CYCLOPHILIN C COMPLEXED WITH CYCLOSPORIN A 2RMC 3	
	2ттс	4	9	171	8.40E-41	0.39	-		COMPLEX (ISOMERASE/IMMUNOSUPP (RESSANT) CYCLOPHILIN C COMPLEXED WITH CYCLOSPORIN A 2RMC 3	
	Isig		219	335	0.0028	0.33	0.01		RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;	TRANSCRIPTION REGULATION SIGMATO, RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION
	1bor		319	363	3.40E-06	-0.43	0.03		TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR

SeqFold Coumpound PDB annotation score	BODIES (PODS), LEUKEMIA, 2 TRÂNSCRIPTION REGULATION	TRANSCRIPTION FACTOR TRANSCRIPTION REGULATION PML: CHAIN: NULL: BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	NEURO-ONCOLOGICAL IMMUNE SYSTEM KH DOMAIN, VENTRAL ANTIGEN I; ALPHA-BETA FOLD, RNA-BINDING CHAIN: A; MOTIF	RNA-BINDING IMMUNE SYSTEM KH DOMAIN, NEUROONCOLOGICAL ALPHA-BETA FOLD RNA-BINDING VENTRAL ANTIGEN 2; MOTIF CHAIN: A, B, C, D;	RNA-BINDING IMMUNE SYSTEM KH DOMAIN, NEUROONCOLOGICAL ALPHA-BETA FOLD RNA-BINDING VENTRAL ANTIGEN 2; MOTIF CHAIN: A, B, C, D;	RNA-BINDING IMMUNE SYSTEM KH DOMAIN, NEUROONCOLOGICAL ALPHA-BETA FOLD RNA-BINDING VENTRAL ANTIGEN 2; MOTIF CHAIN: A, B, C, D;	RNA-BINDING NEUROONCOLOGICAL VENTRAL ANTIGEN 2; CHAIN: A, B, C, D:	RNA-BINDING PROTEIN RNA-BINDING PROTEIN/RNA NOVA-2: CHAIN: A, B; 20- MER RNA HAIRPIN; CHAIN: C, D; MOTHER BINDING PROTEIN; KH DOMAIN, ALPHA-BETA FOLD, RNA-BINDING MOTHER PROTEIN/RNA 2 STRUCTURE	SIGNAL TRANSDUCTION LIGASE CBL, UBCH7, ZAP-70, E2, PROTEIN CBL; CHAIN: A; UBIQUITIN, E3, PHOSPHORYLATION, ZAP-70 PEPTIDE; CHAIN: B; 2 TYROSINE KINASE, IDEICHTEN CONTING ATTACK
PMF		0.16	8.0	0.98	0.95	0.88	0.99	0.99		0.71
Verify score		-0.5	-0.21	0.49	0.42	0.44	0.3	0.51	0.66	0.28
PSI- BLAST		9.80E-09	3.40E-12	2.80E-14	2.80E-14	2.80E-05	4.20E-16	2.80E-15	8.40E-15	3.40E-13
End AA		366	370	104	104	106	104	104	109	370
Start AA		321	321	. 39	39	39	39	39	39	320
Chain ID				<	∀	B	ι O	Ω	«	4
PDB ID		1 bor	1chc	1dt4	1dtj	1dtj	1dtj	1dtj	lec6	1fbv
SEQ		549	549	549	549	549	549	549	549	549

SEQ ID	PDB ID	Chain 1D	Start AA	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
									BINDING SITE; CHAIN: B, C;	
554	lme y	9	1006	1032	1.00E-10	0.14	-0.19		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA
									PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2 CENTER STRIPE COMPLEY
										(ZINC FINGER/DNA)
554	1me	G	443	470	1.70E-12	0.27	0.52		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
	>	·							CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
						_			PROTEIN; CHAIN: C, F, G;	CRYSTAL STRUCTURE COMPLEY
										CRISIAL SIROCIONE, COMPLESA (ZINC FINGER/DNA)
554	Ime	ŋ	096	985	5.10E-09	-0.21	0.04		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
	>								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
							_		FROIEIN; CHAIN: C, F, G,	CRYSTAL STRUCTURE, COMPLEX
										(ZINC FINGER/DNA)
554	<u>F</u>	<	366	470	3.40E-13	-0.03	0.04		TRANSCRIPTION FACTOR	COMPLEX (TRANSCRIPTION
		_							IIIA; CHAIN: A; 55 KNA	KEGULATION/DNA) IFIIIA; 35 GENE;
									GEINE, CHAIN: E, F;	TRANSCRIPTION FACTOR SCIENA 2
			_			_		_		GENE. DNA BINDING PROTEIN, ZINC
										FINGER, COMPLEX 3
										(TRANSCRIPTION REGULATION/DNA)
554	2adr		409	472	5.10E-13	0.04	0.24		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION
										TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR
554	2adr		494	553	1.70E-15	-0.35	0		ADRI; CHAIN: NULL;	TRANSCRIPTION REGULATION
										TRANSCRIPTION REGULATION, ADRI 21NC FINGER NMR
554	2drp	A	401	472	2.80E-11	0.01	0.89		COMPLEX(TRANSCRIPTION	
									REGULATION/DNA)	
									TWO 21NC-FINGER	
									PEPTIDE) COMPLEXED	
									WITH 2DRP 3 DNA 2DRP 4	
555	1	•	30	205	3 40E 16	2			TA ANG CHIRTING 141	The appropriate representation of the state
CCC	161	<	25	293	3.40E-13	-0.15)		IKANSCKIPIIONAL REPRESSOR TUPI; CHAIN: A, B, C;	IRANSCRIPTION INHIBITOR BELA- PROPELLER
555	1 got	В	136	352	1.70E-11	0.11	-0.13		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT-	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1,

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
e Š	<u>e</u>	9	∀	V V	BLAST	score	score	score		
									METHYLTRANSFERASE; CHAIN: A, B, C, D;	METHYLTRANSFERASE
558	1d2h	V	19	223	2.80E-13	-0.17	0.19		GLYCINE N- METHYLTRANSFERASE; CHAIN: A, B, C, D;	TRANSFERASE METHYLTRANSFERASE
558	1dus	4	89	187	1.40E-12	0.35	68.0		MJ0882; CHAIN: A;	STRUCTURAL GENOMICS HYPOTHETICAL PROTEIN, METHANOCOCCUS JANNASCHII
558	1dus	<	83	213	2.80E-14	0.25	0.93		MJ0882; CHAIN: A;	STRUCTURAL GENOMICS HYPOTHETICAL PROTEIN, METHANOCOCCUS JANNASCHII
558	1869	-	43	195	1.70E-26	0.11	0.31		HNRNP ARGININE N- METHYL TRANSFERASE; CHAIN: 1, 2, 3, 4, 5, 6;	TRANSFERASE SAM-BINDING DOMAIN, BETA-BARREL, MIXED ALPHA-BETA, HEXAMER, 2 DIMER
558	lxva	<	43	195	5.10E-21	-0.08	0.72		GLYCINE N- METHYLTRANSFERASE; CHAIN: A, B;	METHYLTRANSFERASE GNMT, S- ADENOSYL-L-METHIONINEN: GLYCINE METHYLTRANSFERASE
558	lyub		71	157	1.20E-06	0.44	90.0		RRNA METHYLTRANSFERASE; CHAIN: NULL;	METHYLTRANSFERASE ERMAM; METHYLTRANSFERASE, ERM, ERMAM, MLS ANTIBIOTICS, NMR, 2 RRNA
558	2ad m	A	62	681	1.20E-08	0.09	0.48	·	ADENINE-N6-DNA- METHYLTRANSFERASE TAQI; CHAIN: A, B;	METHYLTRANSFERASE TRANSFERASE, METHYLTRANSFERASE, RESTRICTION SYSTEM
559	lgwz		225	301	4.20E-05	-0.17	0.25		SHP-1; CHAIN: NULL;	HYDROLASE PROTEIN-TYROSINE PHOSPHATASE; HYDROLASE, PROTEIN TYROSINE PHOSPHATASE, CATALYTIC DOMAIN, 2 WPD LOOP, SH2 DOMAIN
559	P P		158	298	3.40E-39			137.8	PYST1; CHAIN: NULL;	HYDROLASE DUAL SPECIFICITY PHOSPHATASE, MAP KINASE HYDROLASE
559-	1mk P		159	287	3.40E-39	0.88	_		PYSTI; CHAIN: NULL;	HYDROLASE DUAL SPECIFICITY PHOSPHATASE. MAP KINASE HYDROLASE
559	l vhr	Ą	135	307	1.10E-34			96.19	HUMAN VHI-RELATED DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B;	HYDROLASE VHR; HYDROLASE, PROTEIN DUAL-SPECIFICITY PHOSPHATASE
559	1vhr	A	142	298	1.10E-34	0.78			HUMAN VHI-RELATED	HYDROLASE VHR; HYDROLASE,

	Chain ID	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
								DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B;	PROTEIN DUAL-SPECIFICITY PHOSPHATASE
<		146	282	1.00E-32	0.29	_		HUMAN VHI-RELATED DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B,	HYDROLASE VHR; HYDROLASE, PROTEIN DUAL-SPECIFICITY PHOSPHATASE
∀		188	248	6.80E-22	0.4	0.77		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
-	∢	194	276	1.50E-28	0.03	0.83		QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
ļ	¥	194	278	1.50E-28			56.74	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
ļ ·	<	222	288	1.20E-24	-0.45	0.86		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
ļ	၁	187	246	6.80E-36	0.07	0.69		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	ပ	193	276	5.10E-49	0.08			DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	C	193	277	5.10E-49			69.14	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	၁	221	288	3.40E-40	-0.37	0.46		DNA: CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX

PDB Chain Start End PSI- ID ID AA BLAST Ime G 219 246 1.70E-13	Chain Start End ID AA AA 219 246 1.	A AA 246 1.		PSI. BLAS	3	Verify score	PMF score 0.92	SeqFold	Coumpound DNA; CHAIN: A, B, D, E;	PDB annotation (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA) ZINC
y 143 A 193 280 6.80E-18	193	280	-	6 805-18				50 62	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; TRANSCRIPTION FACTOR	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA) COMPLEX (TRANSCRIPTION
007 CC1	661	267		010000				70.00	IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F;	REGULATION AND TELLIA SS GENE; NMR, TELLIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION DNA)
1tß A 194 276 6.80E-18	194 276	276		6.80E-18		-0.15	0.01		TRANSCRIPTION FACTOR IIIA; CHAIN: A, 5S RNA GENE; CHAIN: E. F;	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA, 5S GENE: NMR. TFIIIA, PROTEIN. DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)
1tf6 A 198 277 1.40E-20	198 277	277		1.40E-20		-0.28	0.16		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN
1ubd C 167 277 6.80E-29	167 277	277		6.80E-29				56.56	YY I, CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
1ubd C 186 276 6.80E-29	186 276	276		6.805-29		-0.32	0.89		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
1ubd C 201 284 5.10E-27	201 284	284	Ħ	5.10E-27		-0.02	0.88		YYI; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION

	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation	
									ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	
+	2gli	<	133	278	1.50E-26			57.42	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	
+	2gli	<	188	278	1.50E-26	0.23	0.72		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	
	163u	٧	_	605	1.70E-40		ļ	161.25	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	
	163u	<	6	909	1.70E-40	-0.03	0.71		PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	
 	lee4	∢	144	602	1.30E-43	0.42	-		KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	
 	lee4	∢	18	459	1.40E-31	-0.06	0.83	·	KARYOPHERIN ALPHA: CHAIN: A. B. MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F,	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	
	1ec4	₹	184	109	1.70E-37	0.44	_		KARYOPHERIN ALPHA; CHAIN: A. B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C. D. E. F;	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	
	lec4	⋖	7	390	3.40E-28	0.34			KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	
	lial	×.	304	603	5.10E-21	0.38	96.0		IMPORTIN ALPHA; CHÁIN: A;	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2	

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
Ö										ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION
571	lial	<	33	475	1.40E-32			152.5	IMPORTIN ALPHA; CHAIN: A;	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION
571	lial	<	530	607	1.70E-09	0.26	0.19		IMPORTIN ALPHA; CHAIN: A;	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION
571	lial	<		390	1.40E-32	0.18	0.88		IMPORTIN ALPHA; CHAIN: A;	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION
571	libr	B	225	909	6.80E-13	-0.01	0.4		RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	SMALL GTPASE KARYOPHERIN BETA, P9S SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR
571	libr	В	3	97	5.10E-12	0.11	-0.09		RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE. NUCLEAR TRANSPORT RECEPTOR
571	1qbk	В	9	\$09	3.40E-49	10.0	0.74		KARYOPHERIN BETA2; CHAIN: B; RAN; CHAIN: C;	NUCLEAR TRANSPORT PROTEIN COMPLEX HEAT REPEATS, NUCLEAR TRANSPORT PROTEIN COMPLEX
571	lqgr	<	E.	474	6.80E-23	-0.16	0.16		IMPORTIN BETA SUBUNIT; CHAIN: A; IMPORTIN ALPHA-2 SUBUNIT; CHAIN: B;	TRANSPORT RECEPTOR KARYOPHERIN BETA-1, NUCLEAR FACTOR P97, IMPORTIN IMPORTIN ALPHA-2 SUBUNIT, KARYOPHERIN ALPHA-2 TRANSPORT RECEPTOR, NUCLEAR IMPORT, HEAT MOTIF, NI.S-BINDING
571	2bct		8	200	1.40E-22	0.29	0.64		BETA-CATENIN; CHAIN:	STRUCTURAL PROTEIN ARMADILLO

SEQ	BOA	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
<u> </u>	a -	a	V V	¥	BLASI	score	score	Score		
									NULL;	REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN
571	2bct		89	209	6.80E-44			160.13	BETA-CATENIN; CHAIN: NULL;	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN
571	2bct		6	431	8.40E-25	0.39	-	·	BETA-CATENIN; CHAIN: NULL;	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN
571	2bct		16	607	6.80E-44	0.31	0.94		BETA-CATENIN; CHAIN: NULL;	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN
571	3bct			386	6.80E-18	0.57	-1.41		BETA-CATENIN: CHAIN: NULL:	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
571	3bct		13	476	1.40E-31			141.33	BETA-CATENIN; CHAIN: NULL;	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
571	3bct		138	604	5.10E-36	0.45	1202.08		BETA-CATENIN; CHAIN: NULL;	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
571	3bct		51	430	1.40E-31	0.35	-		BETA-CATENIN; CHAIN: NULL;	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
572	lav!	<	23	219	5.60E-11			67.34	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	LIPID TRANSPORT APO A-I; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION
572	Icun	<	9	219	2.80E-12			62.14	ALPHA SPECTRIN; CHAIN: A, B, C;	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN
572	Iquu	4	£	245	1.40E-13			67.92	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN
572	lsig		31	161	2.80E-09	-0.28	0.02		RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION

				,,,	41; 1P, IN 2 LY,	BCI;	EC1;	35 XI,	IN 2 LY.
PDB annotation	LIPID TRANSPORT APO A-1; LIPOPROTEIN, LIPID TRANSPORT, CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT- ACTIVATION	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COILED-COILS, STRUCTURAL PROTEIN	SIGNALING PROTEIN GUANINE NUCLEOTIDE- BINDING PROTEIN 1; GBP, GTP HYDROLYSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 RELATED, LARGE GTPASE FAMILY, SIGNALING PROTEIN	ENDOCYTOSIS/EXOCYTOSIS NSECI; PROTEIN-PROTEIN COMPLEX. MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS NSEC!, PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	SIGNALING PROTEIN GBP, GTP HYDROLYSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 RELATED, LARGE GTPASE FAMILY. GMPPNP, GPPNHP.
	LIPII LIPO CHO ATH ACTI	COL CHA TRA	STR(REPH HEL) TAN STR(STRI REPI HEL TAN STRI	SIGN NUC GBP INTE	PRO MUI	END PRO MUI	SYN BUND	SIGN HYT INTE GMB
Coumpound	APOLIPOPROTEIN A-1; CHAIN: A, B, C, D;	COLICIN IA; CHAIN: NULL;	A. B. C.	ALPHA SPECTRIN; CHAIN: A, B, C;	INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN 1; CHAIN: A;	SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN BINDING PROTEIN 1, CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN-1A; CHAIN: A, B, C;	INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN I; CHAIN: A;
SeqFold score									·
PMF	-0.18	-0.2	10:0-	-0.12	-0.13	-0.17	-0.19	-0.18	-0.19
Verify	0.52	90:00	0.5	0.57	0.43	0.39	0.31	0.43	0.29
PSI- BLAST	7.00E-13	4.20E-16	5.60E-12	7.00E-15	1.40E-10	8.40E-14	1.40E-11	1.10E-16	4.20E-11
End	137	175	175	125	140	145	175	129	145
Start AA	20	7	-	9	16	4	8	14	11
Chain ID	V V		₹	<	4	В	В	<	∢
PDB ID	lavl	lcii	Icun	lcun	14g3	ldn1	ldn1	lez3	1f5n
SEQ ID NO:	573	573	573	573	573	573	573	573	573

PDB annotation	PROTEIN TRANSPORT HELIX-TURN- HELIX TPR-LIKE REPEAT, PROTEIN TRANSPORT	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TRIPLE. HELIX COILED COIL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TRIPLE- HELIX COILED COIL, CONTRACTILE PROTEIN	COMPLEX (TRANSDUCER/TRANSDUCTION) GT BETA-GAMMA; MEKA, PP33; PHOSDUCIN, TRANSDUCIN, BETA-GAMMA, SIGNAL TRANSDUCTION, 2 GAMMA, SIGNAL TRANSDUCTION, 2 GEQULATION, PHOSPHORYLATION, G PROTEINS, THIOREDOXIN, 3 VISION, MEKA, COMPLEX (TRANSDUCER/TRANSDUCTION)	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- PROPELLER	TRANSCRIPTION INHIBITOR BETA-
Coumpound	VESICULAR TRANSPORT PROTEIN SEC17; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	TRANSDUCIN; CHAIN: B. G; PHOSDUCIN; CHAIN: P;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C.	TRANSCRIPTIONAL
SeqFold score			53.17									
PMF	-0.2	-0.2		-0.18	-0.19	0.22	0.1	0	0.22	-0.12	-0.14	1
Verify score	0.2	0.4		0.36	0.08	0.47	0.37	0.41	-0.14	0.22	0.14	8.0
PSI- BLAST	1.40E-12	5.60E-12	1.40E-19	1.40E-19	8.40E-11	9.80E-18	5.60E-16	4.20E-15	5.10E-05	1.40E-09	5.60E-91	1.70E-76
End	146	101	250	174	144	358	466	484	476	209	484	484
Start	9	_	9	7		110	172	262	353	76	601	193
Chain LD	V	A	<	4	۵.	<	4	<	V	<	V	A
PDB ID	lqqe	Iquu	Iquu	lquu	2trc	lcrz	lcrz	lcrz	lcrz	lcrz	lerj	leri
SEQ ID NO:	573	573	573	573	573	574	574	574	574	574	574	574

PDB annotation	A, PROPELLER	A, PROPELLER	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDICTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDICTE) COMPLEX
Coumpound	REPRESSOR TUP1; CHAIN: A, B, C.	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA CHIMERA, CHAIN: A, GT- BETA; CHAIN: B, GT- GAMMA, CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;
SeqFold score						
PMF		_		-	-	66.0
Verify score		0.46	0.95	0.7	0.79	0.43
PSI- BLAST		1.70E-67	3.40E-71	1.00E-50	6.80E-79	3.40E-55
End		356	400	273	483	315
Start		48	101	15	061	45
Chain ID		Y	В	В	B	В
PDB ID		lerj	1got	lgot	1got	Igot
SEQ TO NO:		574	574	574	574	574

PDB annotation	SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	SE, HYDROLASE PROLYL ENDOPEPTIDASE, POST-PROLINE CLEAVING PROLYL OLIGOPEPTIDASE, AMNESIA, ALPHA/BETA-HYDROLASE, BETA- 2 PROPELLER	ITE OXIDOREDUCTASE ENZYME, B; NITRITE REDUCTASE, OXIDOREDUCTASE, DENITRIFICATION, 2 ELECTRON TRANSPORT, PERIPLASMIC	L; TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2 PROTEIN	-\$d>	VPS-	-\$dp	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF IL-6 TYPE CYTOKINES, THIRD 2 N-TERMINAL DOMAIN. TRANSMEMBRANE, GLYCOPROTEIN	T-CELL SURFACE GLYCOPROTEIN
Coumpound		PROLYL OLIGOPEPTIDASE; CHAIN: A;	CYTOCHROME CDI NITRITE REDUCTASE; CHAIN: A, B;	COLICIN IA; CHAIN: NULL;	DEFENSIN DEFENSIN /HNP\$-3 1DFN 3	DEFENSIN DEFENSIN /HNP\$-3 1DFN 3	DEFENSIN DEFENSIN /HNP\$.	HEMOLIN; CHAIN: A, B;	HEMOLIN; CHAIN: A, B;	GP130; CHAIN: NULL;	T-CELL SURFACE
SeqFold score				98.21	59.92			111.17			
PMF score		0.05	-0.19			_	-		0.05	-0.08	-0.01
Verify		-0.33	0.43			-0.35	-0.35		0	0.22	0.43
PSI- BLAST		0.00056	5.60E-79	1.705-10	1.10E-12	1.10E-12	5.10E-11	1.70E-47	1.70E-47	5.60E-11	1.40E-19
End		172	478	648	101	101	101	435	416	434	220
Start AA		=	104	49	72	73	73	30	32	344	49
Chain ID		<	<		V	V	Ą	<	4		
PDB ID		1qfm	19ks	1cii	1dfn	1dfn	ıJp!	1bih	l bih	1bj8	Icdy
SEQ NO:		574	574	575	577	577	577	578	578	578	578

PDB annotation	TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC, LIPOPROTEIN, T- CELL SURFACE GLYCOPROTEIN		CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	IMMUNE SYSTEM ABZYME TRANSITION STATE ANALOG, IMMUNE SYSTEM	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN
Coumpound	CHAIN: NULL;	NEURAL ADHESION MOLECULE DROSOPHILA NEUROGLIAN (CHYMOTRYPTIC FRAGMENT CONTAINING THE ICFB 3 TWO AMINO PROXIMAL FIBRONECTIN TYPE III REPEATS ICFB 4 (RESIDUES 610 - 814)) ICFB 5	AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	7C8 FAB FRAGMENT; SHORT CHAIN; CHAIN; A, C; 7C8 FAB FRAGMENT; LONG CHAIN; CHAIN; B, D	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	NEURAL CELL ADHESION MOLECULE: CHAIN: A, B, C, D;
SeqFold score									
PMF score		0.39	-0.11	0	-0.15	-1.41	0.43	0.34	0.99
Verify		0.25	60:0	90.0	0.11	0.1	0.18	1.0	0.16
PSI- BLAST		7.00E-20	1.40E-48	1.40E-50	6.80E-13	1.40E-44	3.40E-28	1,40E-30	5.60E-30
End		435	522	436	328	340	225	220	214
Start		252	137	32	140	134	31	36	33
Chain ID			<	A	¥	ن ا	D	Q	A
PDB ID		lcfb	1cs6	lcs6	1ct8		lcvs	lcvs	lepf
SEQ ID		578	878	578	578	578	578	578	578

	FACTOR E HE I-SET E	I FACTOR E HE I-SET E	I FACTOR LE 'HE I-SET 'E	TOR 4-HELICAL EX, FN 2 DOMAINS, PLEX	зета	KGD, 1FNF 18	NDING				
PDB annotation	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE 1-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1; FGFR1; IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	HORMONE/GROWTH FACTOR/HORMONE RECEPTOR 4- HELICAL BUNDLE, ALPHA HELICAL BUNDLE, TERNARY COMPLEX, FN 2 III DOMAINS, BETA SHEET DOMAINS, CYTOKINE-RECEPTOR COMPLEX	CONTRACTILE PROTEIN IMMUNOGLOBULIN FOLD, BETA BARREL	CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX IFNF 18	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	-			
Coumpound	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR I; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR I; CHAIN: C, D;	PLÁCENTAL LACTOGEN; CHAIN: A: PROLACTIN RECEPTOR: CHAIN: B, C;	TELOKIN; CHAIN: A	FIBRONECTIN; IFNF 6 CHAIN: NULL; IFNF 7	FIBRONECTIN; CHAIN: A;	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) 1HNG 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (RAT) 14NG 3	
SeqFold score											
PMF score	0.25	0.48	0.01	0.17	-0.05	0.46	0.04	0.29	-0.12	0.15	
Verify score	60.0	90.0-	0.03	0.29	90.0	0.24	0.24	0.3	0.05	0.15	
PSI- BLAST	4.20E-30	2.80E-31	5.60E-28	1.40E-11	6.80E-18	5.60E-15	1.40E-12	1.40E-18	4.20E-14	2.80E-24	
End AA	220	228	220	436	131	432	432	211	292	227	
Start	38	36	29	267	27	264	264	47	143	49	
Chain ID	യ	Ð	U	B	∢		٧		٧	A	
PDB ID	lev2	lev2	levt	166	Ifhg	Juji	1 fnh	lhnf	lhng	Ihng	
SEQ NO.	578	578	578	578	578	878	878	578	578	578	

PDB annotation	IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION	COMPLEX (IMMUNOGLOBULIN/RECEPTOR) IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, RECEPTOR, 2 SIGNAL, COMPLEX (IMMUNOGLOBULIN/RECEPTOR)		CELL ADHESION PROTEIN CELL ADHESION PROTEIN, RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTMS; CELL ADHESION, GLYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTM5; CELL ADHESION, GLYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTMS; CELL ADHESION, GLYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN
Coumpound	MAB61.1.3; CHAIN: A, B, C, D	INTERLEUKIN-I BETA; CHAIN: A; TYPE I INTERLEUKIN-I RECEPTOR; CHAIN: B;	IMMUNOGLOBULIN IMMUNOGLOBULIN GI (IGGI) (MCG) WITH A HINGE DELETION IMCO 3	FIBRONECTIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	TITIN; CHAIN: NULL;	NTEGRIN BETA-4 SÜBUNIT; CHAIN: A, B;
SeqFold score			96.36					
PMF		-0.12		0.33	0.88	-0.06	-0.13	0.52
Verify score		0.03		0.39	0.29	0.06	0.13	0.3
PSI- BLAST		6.80E-34	3.40E-10	1.30E-17	1.40E-18	4.20E-15	6.80E-15	1.40E-20
End AA		338	423	432	227	132	132	450
Start		22	32	264	139	31	31	251
Chain ID		В	H					<
PDB TD		lifb	lmc o	1mfn	Inct	Inct	Inct	1գ <u>ც</u> 3
SEQ D NO:		578	578	578	578	578	578	578

SEQ ID NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
578	1914	<	264	435	2.80E-17	0.52	0.82		TENASCIN; CHAIN: A, B;	STRUCTURAL PROTEIN TENASCIN, FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN
	1tnm		139	722	7.00E-18	0.39	0.89		MUSCLE PROTEIN TITIN MODULE MS (CONNECTIN) 1TNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) 1TNM 4 1TNM 58	
T	Itam		33	132	8.40E-15	-0.04	0.03		MUSCLE PROTEIN TITTN MODULE MS (CONNECTIN) ITNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) ITNM 4 ITNM 58	
578	lttf		346	434	1.40E-10	0.2	0.05		GLYCOPROTEIN FIBRONECTIN (TENTH TYPE III MODULE) (NMR, 36 STRUCTURES) 1TTF 3	
578	lwio	∢	49	281	2.80E-20	0.04	-0.12		T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	GLYCOPROTEIN CD4; IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM
578	1 wit		139	226	2.80E-17	0.52	0.78		TWITCHIN 18TH 1GSF MODULE; CHAIN: NULL;	MUSCLE PROTEIN IMMUNOGLOBULIN SUPERFAMILY, I SET, MUSCLE PROTEIN
	l wit		31	132	2.80E-15	0.23	0.1		TWITCHIN 18TH IGSF MODULE, CHAIN: NULL,	MUSCLE PROTEIN IMMUNOGLOBULIN SUPERFAMILY, I SET, MUSCLE PROTEIN
578	lww w	×	142	228	5.60E-18	0.02	-0.01		NERVE GROWTH FACTOR; CHAIN: V, W; TRKA RECEPTOR; CHAIN: X, Y;	NERVE GROWTH FACTOR/TRKA COMPLEX BETA-NGF; COMPLEX, TRKA RECEPTOR, NERVE GROWTH FACTOR, CYSTEINE KNOT, 2 IMMUNOGLOBULIN LIKE DOMAIN, NERVE GROWTH FACTOR/TRKA COMPLEX
578	lzxq		31	228	1.10E-30	0.18	0.23		INTERCELLULAR ADHESION MOLECULE-2; CHAIN: NULL;	CELL ADHESION ICAM-2; IMMUNOGLOBULIN FOLD, CELL ADHESION, GLYCOPROTEIN, 2 TRANSMEMBRANE, REPEAT, SIGNAL

PDB annotation	IMMUNE SYSTEM PS8 NATURAL KILLER CELL RECEPTOR; KIR, NATURAL KILLER RECEPTOR, INHIBITORY RECEPTOR, 2 IMMUNOGLOBULIN	IMMUNE SYSTEM CD32, RECEPTOR, FC, CD32, IMMUNE SYSTEM	PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	COAGULATION FACTOR	CELL ADHESION NCAM DOMAIN 1; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, SIGNAL		CELL ADHESION PROTEIN NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	CELL ADHESION PROTEIN NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN-BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	
Coumpound	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	FC GAMMA RIIB; CHAIN: A;	FIBRONECTIN; CHAIN: A;	HUMAN TISSUE FACTOR; 2HFT 4 CHAIN: NULL; 2HFT 5	NEURAL CELL ADHESION MOLECULE; CHAIN: NULL;	HORMONE/RECEPTOR HUMAN GROWTH HORMONE COMPLEXED WITH ITS RECEPTOR 3HHR 3 (EXTRACELLULAR DOMAIN) 3HHR 4	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1
SeqFold									
PMF	-0.02	99.0	-0.12	-0.08	0.98	-0.09	0.31	0.06	0.01
Verify score	0.25	0.1	0.45	0.13	0.64	90.00	90.0	0.04	-0.56
PSI- BLAST	2.80E-25	2.80E-33	1.40E-12	2.80E-12	5.60E-18	5.60E-17	4.20E-18	7.00E-17	1.00E-09
End	220	228	435	436	227	436	220	134	103
Start	30	31	343	264	139	264	139	32	[9]
Chain ID	<	V	A			В	Ą	∢	
PDB ID	2dli	2fcb	2£nb	2hft	Znc ın	3hhr	3nc m	3nc m	lchc
SEQ ID NO:	578	578	578	578	578	578	578	578	6/5

PDB annotation		TRANSCRIPTION FACTOR BTF2 P44 SUBUNIT, BASIC TRANSCRIPTION FACTOR, ZINC BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J	RECOMBINATION ACTIVATING	PROTEIN I; RAGI, V(D)J	RECOMBINATION, ANTIBODY, MAD,	RING FINGER, 2 ZINC BINUCLEAR	CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J	RECOMBINATION ACTIVATING	PROTEIN 1; RAG1, V(D)J	RECOMBINATION, ANTIBODY, MAD,	RING FINGER, 2 ZINC BINUCLEAR	CLUSTER, ZINC FINGER, DNA.	BINDING PROTEIN	DNA-BINDING PROTEIN V(D)J	RECOMBINATION ACTIVATING	PROTEIN 1; RAG1, V(D)J	RECOMBINATION, ANTIBODY, MAD,	RING FINGER, 2 ZINC BINUCLEAR	CLUSTER, ZINC FINGER, DNA-	BINDING PROTEIN	CALCIUM-BINDING PROTEIN CALB;	CALCIUM++/PHOSPHOLIFID BINDING	PROTEIN, 2 CALCIOM-BINDING	ENDOCYTOSIS/EXOCYTOSIS	SYNAPTOTAGMIN, C2-DOMAIN,	EXOCYTOSIS, NEUROTRANSMITTER	2 RELEASE,	ENDOCYTOSIS/EXOCYTOSIS	HYDROLASE CPLA2;	PHOSPHOLIPASE, LIPID-BINDING,	HYDROLASE	TRANSFERASE CALCIUM++,	PHOSPHOLIPID BINDING PROTEIN,	CALCIUM-DINDING 2 FAUTEIIN
Coumpound	STRUCTURE) 1CHC 4	TFIIH P44 SÜBÜNIT; CHAIN: A;	RAGI: CHAIN: NULL:						RAGI; CHAIN: NULL;							RAGI; CHAIN: NULL;							PROTEIN KINASE C (BETA);	CHAIN: A, B;		SYNAPTOTAGMIN I: CHAIN:	A;		-		CYTOSOLIC	PHOSPHOLIPASE A2; CHAIN:	A, B;	PROTEIN KINASE C, ALPHA	TYPE; CHAIN: A;	
SeqFold score																																				
PMF		0.19	0.33						0.29							0.13							0.94			0.52					0.13			86.0		
Verify score		0	-0.01						-0.78							0.01							0.83			90.0					0.05			0.61		
PSI- BLAST		15000:0	0.0037						0.0007							3.40E-07							1.70E-30			1.70E-23					1.70E-15			3.40E-31		
End		102	268						92							128							1443			1442	!				1444			1443		
Start		15	223						59							63							1329			1325	!				1342			1329		
Chain ID		4																					∀			<					٧			V		
PDB LD		1.00 E+53	lrmd						lrmd						_	1md							1825			1byn					1cjy			1dsy		
SEQ NO:		579	579						579					_		579							280			580					280			280		

Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	VΥ	ΑA	BLAST	score	score	score		
<u> </u>								PHOSPHATIDYLSERINE, PROTEIN KINASE C
2	214	1175	0	0.15	1202.08		PHOSPHATIDYLINOSITOL 3- KINASE CATALYTIC SUBUNIT, CHAIN: A;	PHOSPHOINOSITIDE 3-KINASE GAMMA PTDINS-3-KINASE P110, P13K, P1 3K; PHOSPHOINOSITIDE 3-KINASE
								GAMMA, SECONDARY MESSENGER 2 GENERATION, P13K, P1 3K, WORTMANNIN
2	297	1175	0	0.15	-1.41		PHOSPHATIDYLMOSITOL 3- KINASE CATALYTIC	PHOSPHOINOSITIDE 3-KINASE GAMMA PTDINS-3-KINASE PI10, PI3K;
							SUBUNII; CHAIN: A;	PHOSPHOINOSI I IDE 3-KINASE GAMMA, SECONDARY MESSENGER 2 GENERATION, PI3K, PI 3K
	1342	1444	1.70E-15	-0.02	0.64		PHOSPHOLIPASE A2; CHAIN: NULL;	HYDROLASE CALB DOMAIN: HYDROLASE, C2 DOMAIN, CALB DOMAIN
	1321	1442	8.50E-24	90.0	0.59		CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	
	1328	1441	5.10E-29	0.1	0.39		RABPHILIN 3-A; CHAIN: A;	ENDOCYTOSIS/EXOCYTOSIS C2- DOMAINS, C2B-DOMAIN, RABPHILIN, ENDOCYTOSIS/EXOCYTOSIS
	9	577	0	0.27	-0.2		INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN I; CHAIN: A;	SIGNALING PROTEIN GUANINE NUCLEOTIDE- BINDING PROTEIN 1; GBP, GTP HYDROLYSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 REI ATEN 1 ARGE GTPASE FAMILY
								SIGNALING PROTEIN
	7	577	0	0.47	-0.2		INTERFERON-INDUCED GUANYLATE-BINDING PROTEIN I; CHAIN: A;	SIGNALING PROTEIN GBP, GTP HYDROL YSIS, GDP, GMP, INTERFERON INDUCED, DYNAMIN 2 RELATED, LARGE GTPASE FAMILY. CANDAN.
-								OMITAL, OFFICE
	46	604	0	0.29	-0.2		ALPHA-D-GLUCOSE-1,6- BISPHOSPHATE; 3PMG 4 CHAIN: A. B. 3PMG 5	PHOSPHOTRANSFERASE PHOSPHOGLUCOMUTASE: 3PMG 6 PHOSPHOGLUCOMUTASE 3PMG 13
	53	611	0			129.26	ALPHA-D-GLUCOSE-1,6- BISPHOSPHATE; 3PMG 4	PHOSPHOTRANSFERASE PHOSPHOGLUCOMUTASE; 3PMG 6

PDB annotation	PHOSPHOGLUCOMUTASE 3PMG 13	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION		LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	LIM DOMAIN CONTAINING PROTEINS LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN, ZINC 2 FINGER	LIM DOMAIN CONTAINING PROTEINS LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN, ZINC 2 FINGER	CONTRACTILE LIM DOMAIN, CRP, NMR, MUSCLE DIFFERENTIATION, CONTRACTILE	METAL-BINDING PROTEIN LIM DOMAIN CONTAINING PROTEINS ICTL 15
Coumpound	CHAIN: A, B; 3PMG 5	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, I STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION PROTEIN CBL: CHAIN: A: ZAP-70 PEPTIDE: CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A;	QCRP2 (LIM1); CHAIN: NULL;	QCRP2 (LIM1); CHAIN: NULL;	CRP1; CHAIN: A;	AVIAN CYSTEINE RICH PROTEIN; ICTL 3
SeqFold score											
PMF		0.25	0.11	0.98	6.0	0.48	60.0	0.62	0	0.18	0.16
Verify score		-0.73	-0.62	0.24	0.25	0.19	-0.44	-0.16	-0.1	-0.11	-0.11
PSI- BLAST		8.50E-06	8.40E-08	8.50E-15	1.70E-10	8.40E-08	1.20E-05	9.80E-17	1.40E-11	4.20E-11	1.10E-12
End		59	95	09	99	64	72	592	662	629	659
Start AA		91	17	16	81	17	81	536	597	597	969
Chain ID					<	A	₹			<	
PDB ID		Ibor	1bor	Ichc	1fbv	1g25	1825	la7i	1a7i	1881	lot l
SEQ ID		587	587	587	587	587	587	592	592	592	592

					()	-i.L				, 2									
PDB annotation	SIGNALING PROTEIN LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN	SIGNALING PROTEIN LIM DOMAIN CONTAINING PROTEINS, METAL-BINDING PROTEIN	METAL-BINDING PROTEIN CRIP; METAL-BINDING PROTEIN, LIM DOMAIN PROTEIN	METAL-BINDING PROTEIN CRIP, METAL-BINDING PROTEIN, LIM DOMAIN PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ANTI-ONCOGENE CELL CYCLE, ANTI- ONCOGENE, REPEAT, ANK REPEAT	COMPLEX (TRANSCRIPTION	REGULATION/DNA) GABPALPHA; GABPBETA1: COMPLEX	(TRANSCRIPTION	REGULATION/DNA), DNA-BINDING, 2	ANK YRIN REPEATS, TRANSCRIPTION	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANX YRN MOTTE	COMPLEX (KINASE/ANTI-	ONCOGENE) CDK6; P16INK4A, MTS1;	CYCLIN DEPENDENT MINASE, CYCLIN DEPENDENT KINASE	INHIBITORY 2 PROTEIN, CDK, INK4,	CELL CYCLE, MULTIPLE TUMOR	SUPPRESSOR, 3 MTSI, COMPLEX (KINASE/ANTI-ONCOGENE) HEADER	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR
Coumpound	CYSTEINE AND GLYCINE- RICH PROTEIN CRP2; CHAIN: A;	CYSTEINE AND GLYCINE- RICH PROTEIN CRP2; CHAIN: A;	CYSTEINE RICH INTESTINAL PROTEIN; CHAIN: NULL;	CYSTEINE RICH INTESTINAL PROTEIN; CHAIN: NULL;	DNA, CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G,	TUMOR SUPPRESSOR P16INK4A; CHAIN: NULL;	GA BINDING PROTEIN	ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1:	CHAIN: B; DNA; CHAIN: D, E;			PI9INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;	MULTIPLE LUMOR STIPPRESSOR · CHAIN B:	(a :: III III (a :: III III)			CYCLIN-DEPENDENT KINASE 6; CHAIN: A;
SeqFold score																			
PMF	8.0	0.23	0.29	0.07	-0.19	0.72	0.1					0.76	0.82						0.84
Verify score	0.43	-0.27	-0.42	0.15	0.16	0.1	0.35					0.28	0.02						0.23
PSI- BLAST	1.40E-16	1.10E-13	1.30E-19	4.20E-13	1.20E-10	0.00014	0.00014					0.00011	9.80E-05						0.00011
End	592	661	609	663	395	370	369					370	367						367
Start	535	595	537	965	367	315	315					315	315						315
Chain ID	V	4			Ð		В						В						В
PDB ID	lcxx	lcxx	Lim	lmi L	Jme y	lase	lawc					1bd8	1bi7						1blx
SEQ ID NO:	592	592	592	592	592	593	593					593	593						593

~	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
 A Ş	<u>e</u>	<u>a</u>	VV	¥¥	BLAST	score	score	score		
									PI9INK4D; CHAIN: B;	PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
593	191×	В	315	370	0.00011	0.42	0.76		CYCLIN-DEPENDENT KINASE 6: CHAIN: A; PI9INK4D; CHAIN: B;	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT
										KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
593	1498	4	315	370	1.40E-05	-0.02	0.21		CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT
593	1dcq	V	315	370	8.40E-05	0.4	0.76		PYK2-ASSOCIATED PROTEIN BETA; CHAIN: A;	METAL BINDING PROTEIN ZINC- BINDING MODULE, ANK YRIN REPEATS, METAL BINDING PROTEIN
593	ii.	Q	315	370	0.00014	0.31	86:0		NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX
593	Infi	ъ	315	381	2.80E-06	0.25	0.59		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX
594	1a06		19	302	5.10E-90	0.15	_		CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE: CHAIN: NULL;	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN
594	lap 	ம	25	3.8	0	0.23	_		TRANSFERASE(PHOSPHOTR ANSFERASE) \$C-fAMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT)	
									ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (/S1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8	

PDB annotation			PROTEIN KINASE CDK2, PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION, MITOSIS, INHIBITION	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)			
Coumpound	1APM 6	TRANSFERASE (PHOSPHOTR ANSFERASE) \$C-/AMP\$-DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6, CHAIN: A; PI9INK4D; CHAIN: B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	PHOSPHOTRÁNSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) ICTP 3 (CATALYTIC
SeqFold		125.65	120.88	127.11	127.4		127.61
PMF						-	
Verify score						0.33	
PSI- BLAST		0	1.40E-60	7.00E-56	0	0	0
End		325	309	325	328	318	328
Start		n	27	52	-	25	2
Chain ID		<u>u</u>		<	ம	Э	ш
PDB ID		ge E	laq1	1blx	L Cm	lcm k	lctp
SEQ ID NO:	:	594	594	594	594	594	594

PDB annotation					ENDOCYTOSIS/EXOCYTOSIS NSEC1;	PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSIS	SYNAPTOTAGMIN ASSOCIATED 33 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSIS	SYNAPTOTAGMIN ASSOCIATED 35	KDA PROTEIN, P35A, THREE HELIX BUNDLE	TRANSFERASE KINASE DOMAIN,	AUTOINHIBITORY FRAGMENT,	HOMODIMER			TRANSFERASE KINASE DOMAIN,	AUTOINHIBITORY FRAGMENT,	Name of the state		PHOSPHOTRANSFERASE FGFRIK,	FIBROBLAST GROWTH FACTOR	RECEPTOR 1; TRANSFERASE,	TYROSINE-PROTEIN KINASE, ATP-	BECEPTOR PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGFRIK.	FIBROBLAST GROWTH FACTOR	RECEPTOR 1; TRANSFERASE,	TYROSINE-PROTEIN KINASE, ATP-	BINDING, 2 PHOSPHORYLATION,
Coumpound	SUBUNIT) ICTP 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN	KINASE (E.C.2.7.1.37) (CAPK)	ICTP 3 (CATALYTIC	SYNTAXIN BINDING	PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B;	SYNTAXIN-1A; CHAIN: A, B,	ij	SYNTAXIN-1A; CHAIN: A, B,	Ü		SERINE/THREONINE-	PROTEIN KINASE PAK-	ALPHA; CHAIN: A, B;	SERINE/THREONINE-	ALPHA; CHAIN: C, D;	SERINE/THREONINE-	PROTEIN KINASE PAK-	SERINE/THREONINE	PROTEIN KINASE PAK-	FGF RECEPTOR 1; CHAIN: A,	B;				FGF RECEPTOR 1: CHAIN: A.	B;			
SeqFold score																					125.79					130.09				
PMF		_			-0.11		-0.19		-0.18			-					-	1202.08												
Verify score		0.25			90.0		0.1		0.16			0.5					0.43													
PSI- BLAST		0			4.20E-12		1.40E-08		2.80E-09			5.60E-86					1.20E-67				5.10E-34					1.20E-40				
End AA		318			009		592		617			302					300				287					286				
Start		25			442		452		486	_		14					4				20					=		_		
Chain ID		3			В		V		٧			J					S				A					В				
PDB D		1ctp			Idh.		1623		lez3			113m					113m				Ifgk					l fgk)			
SEQ NO:		594			594		594		594			594					594				594					594				

Coumpound PDB annotation	RECEPTOR, PHOSPHOTRANSFERASE	HUMAN CYCLIN- DEPENDENT KINASE 2; TRANSFERASE, SERINE/THREONINE CHAIN: NULL; CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	HUMAN CYCLIN- DEPENDENT KINASE 2; TRANSFERASE, SERINE/THREONINE CHAIN: NULL; CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	TWITCHIN; CHAIN: NULL; KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TWITCHIN; CHAIN: A, B; KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	MAP KINASE P38; CHAIN: TRANSFERASE MITOGEN NULL; TRANSFERASE, MAP KINASE, SERINE/THREONINE-PROTEIN KINASE, 2 P38	PHOSPHORYLASE KINASE; KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE. PROTEIN, 2 KINASE, ATP-BINDING, CALMODULIN-BINDING	PHOSPHORYLASE KINASE; KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE-PROTEIN, 2 KINASE, ATP-BINDING, CALMODULIN-BINDING	EXTRACELLULAR REGULATED KINASE 2; ACTIVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, ERK2; TRANSFERASE, SERINE/THREONINE-PROTEIN KINASE, MAP KINASE, 2 ERK2	
PI a		HUMAN CYCL DEPENDENT K CHAIN: NULL;	HUMAN CYCL DEPENDENT K CHAIN: NULL;	TWITCE	TWITCH			PHOSPI		
SegFold			137.2			130.84	127.79		122.57	_
PMF		_		_	-					
Verify score		0.48		0.5	0.48			0.53		
PSI- BLAST		6.80E-62	6.80E-62	3.40E-71	1.70E-71	1.20E-51	6.80E-88	6.80E-88	8.50E-50	
End AA		284	312	285	284	356	285	282	325	
Start		25	27	20	20	9	61	24	10	
Chain ID					<					
PDB ID		Ihcl	The	Ikoa	1kob	1p38	1phk	1 phk	3erk	_
SEQ ID		594	594	594	594	594	594	594	594	

PSI- Verify BLAST score 4.20E-07 -0.44 3.40E-100 -0.03
8.50E-49 0.03
8.50E-32 -0.06
1.70E-35 0.2
3.40E-30 0.01
5.60E-09 -0.08

PDB annotation	KINASE 1CKI 18	TRANSFERASE STRESS-ACTIVATED PROTEIN KINASE-3, ERK6, ERK5: P38-GAMMA, GAMMA, PHOSPHORYLATION, MAP KINASE	•		TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	PHOSPHOTRANSFERASE FGFRIK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP-BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGFRIK, FIBROBLAST GROWTH FACTOR RECEPTOR I; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATPBINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	TRANSFERASE P150, C-ABL; KINASE, KINASE INHIBITOR, STI-571, ACTIVATION LOOP	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION,
Coumpound	ICKI 6 CHAIN: A, B; ICKI 7	PHOSPHORYLATED MAP KINASE P38-GAMMA; CHAIN: A, B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) ICTP 3 (CATALYTIC SUBUNIT) ICTP 4	SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: C, D;	FGF RECEPTOR 1; CHAIN: A, B;	FGF RECEPTOR 1; CHAIN: A, B;	PROTO-ONCOGENE TYROSINE-PROTEIN KINASE ABL; CHAIN: A, B;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;
SeqFold score									
PMF score		60.0	0.37	0.48	0.89	0.22	0.33	0.51	0.7
Verify score		0.32	-0.11	0.18	0.35	0.14	-0.07	0.12	0.26
PSI- BLAST		1.50E-31	0_	6.80E-97	6.80E-46	5.10E-31	7.00E-08	4.20E-06	1.70E-47
End		396	434	434	397	344	372	372	399
Start AA		163	122	122	148	147	206	192	146
Chain ID		A	வ	ங	ပ	В	æ	V	
PDB ID		lcm 8	r LCm	1ctp	1f3m	1fgk	1fgk	1fpu	1hcl
SEQ ID		969	965	965	969	969	969	296	596

PDB annotation	MITOSIS, PHOSPHORYLATION	SERINE/THREONINE-PROTEIN KINASE CSBP, RK, P38, PROTEIN	SER/THR-KINASE, SERINE/THREONINE-PROTEIN KINASE	TRANSFERASE INK3; TRANSFERASE,	INK3 MAP KINASE, SERINE/THREONINE PROTEIN 2	KINASE	TRANSFERASE JNK3: TRANSFERASE, INK3 MAP KINASE	SERINE/THREONINE PROTEIN 2 KINASE	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE	TRANSFERASE, MAP KINASE,	SERINE/THREONINE-PROTEIN KINASE, 2 P38	 -	PHOSPHORYLASE KINASE;	TRANSFERASE, SERINE/THREONINE-	PROTEIN, 2 KINASE, ATP-BINDING,	SERINE KINASE SERINE KINASE,	TRANSFERASE AAC; AMINOGLYCOSIDE 6'-N-	ACETYLTRANSFERASE, ANTIBIOTIC	Z KESISTANCE, ACETTE COENZIME A		(A; HYDROLASE ERA, GTPASE, RNA-BINDING, RAS-LIKE, HYDROLASE	
Coumpound		P38 MAP KINASE; CHAIN: NULL;		C-JUN N-TERMINAL	KINASE; CHAIN: NULL;		C-IUN N-TERMINAL		TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN:	,,,,,,		PHOSPHORYLASE KINASE;	CHAIN: NULL;			TITIN; CHAIN: A, B;	AMINOGLYCOSIDE N6- ACETYLTRANSFERASE	TYPE 1; CHAIN: A;			GTP-BINDING PROTEIN ERA; CHAIN: A, B;	
SeqFold score																								
PMF		0.65		0.74			0.51		99.0	0.76	0.55			0.88				0.92	98.0				0.07	
Verify score		0.1		0			-0.12		0.15	0.31	60:0			0.22				91.0	0.38			-	-0.81	_
PSI- BLAST		1.40E-32		3.40E-34			7.00E-07		1.70E-53	6.80E-54	3.40E-36			1.70E-60				3.40E-41	0.0065				0.00039	
End		349		346			351		400	398	349			397		_		397	145				171	_
Start		147	<u>.</u>	146			258		146	141	147			147				143	17				146	
Chain ID						!				¥								V	¥				4	
PDB ID		lian		ljnk			1 Jnk		Ikoa	1kob	1p38			1phk				1tki	1587				lega	_
SEQ ID NO:		965		965			969		969	969	969			596				969	601				602	_

PDB annotation	N: A, B, ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ittol 3- Phosphotransferase rhogap bomain; phosphotransferase, TPASE ACTIVATING PROTEIN, GAP, CDC42, 2 PHOSPHOINOSITIDE 3- KINASE. SH3 DOMAIN, SH2 DOMAIN, 3 SIGNAL TRANSDUCTION	SITOL 3- DOMAIN; PHOSPHOTRANSFERASE, TPASE ACTIVATING PROTEIN, GAP, CDC42, 2 PHOSPHOINOSITIDE 3- KINASE, SH3 DOMAIN, SH2 DOMAIN, 3 SIGNAL TRANSDUCTION			TL; G-PROTEIN CDC42 GTPASE- ACTIVATING PROTEIN; G-PROTEIN, GAP, SIGNAL-TRANSDUCTION	P. 4; COMPLEX(GTPASE ACTIVATN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP; COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	
Coumpound	SYNTAXIN-1A; CHAIN: A, B, C;	PHOSPHATIDYLINOSITOL 3- KINASE; CHAIN: A, B;	PHOSPHATIDYLMOSITOL 3- KINASE; CHAM: A, B;	PHOSPHATIDYLINOSITOL 3- KINASE; CHAIN: A, B;	RHOGAP; CHAIN: NULL;	RHOGAP; CHAIN: NULL;	P50-RHOGAP; CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;	P50-RHOGAP, CHAIN: A, TRANSFORMING PROTEIN RHOA; CHAIN: B;
SeqFold score					112.2		118.67	
PMF score	0.07	_	_	_		-		_
Verify score	-0.14	0.43	0.78	7.0		0.58		0.54
PSI- BLAST	0.0039	1.30E-20	1.30E-20	2.60E-40	1.80E-30	1.80E-30	9.10E-48	1.10E-30
End	669	192	192	217	212	189	212	201
Start AA	260	25	25	25	13	16	15	16
Chain ID	<	<	æ	æ			∢	<
PDB ID	lez3	Jpb	lpb	dq ≫	lrgp	lrgp	اير 1	1524
SEQ ID NO:	\$09	909	909	\$09	\$09	909	909	909

	PDB ID	Chain 1D	Start AA	End AA	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
1									TRANSFORMING PROTEIN RHOA; CHAIN: B;	ACTIVATN/PROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP, COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP
1	lafi		143	203	0.00026	0.86			MERP; CHAIN: NULL;	MERCURY DETOXIFICATION MERCURIC TRANSPORT PROTEIN: MERCURY DETOXIFICATION, PERJPLASMIC, HEAVY METAL SANDWICH
	law0		143	203	0.0012	0.81	_		MENKES COPPER- TRANSPORTING ATPASE; CHAIN: NULL;	HYDROLASE COPPER- TRANSPORTING ATPASE, COPPER- BINDING DOMAIN, HYDROLASE
	lial	V	21	16	1.20E-06	-0.02	0.46		IMPORTIN ALPHA; CHAIN: A;	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION
	3bct		16	148	0.00026	0.42	96.0		BETA-CATENIN; CHAIN: NULL;	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
	1a09	<	150	255	6.50E-41			172.37	C-SRC TYROSINE KINASE; CHAIN: A, B; ACE-FORMYL PHOSPHOTYR-GLU-(N, N- DIPENTYL AMINE); CHAIN: C, D;	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE)
	1a09	<	150	255	6.50E-41	1.07	-		C-SRC TYROSINE KINASE; CHAIN: A, B; ACE-FORMYL PHOSPHOTYR-GLU-(N, N- DIPENTYL AMINE); CHAIN: C, D;	COMPLEX (TRANSFERASE/PEPTIDE) COMPLEX (TRANSFERASE/PEPTIDE)
	1a81	⋖	126	255	7.80E-33	0.57	_		SYK KINASE; CHAIN: A, C, E, G, I, K; T-CELL SURFACE GLYCOPROTEIN CD3 EPSILON CHAIN; CHAIN: B, D, F, H, J, L;	COMPLEX (TRANSFERASEPEPTIDE) ITAM PEPTIDE; COMPLEX (TRANSFERASE/PEPTIDE), SYK, KINASE, SH2 DOMAIN, ITAM

PDB annotation	D, F, H; KINASE RECEPTOR R4; COMPLEX (ISOMERASE/PROTEIN KINASE), RECEPTOR 2 SERINE/THREONINE KINASE			DSINE V-SRC SH2 DOMAIN SRC SH2; V-SRC SRMING SH2 DOMAIN, PHOSPHOTYROSINE SH2 DOMAIN, PP60 2 SRC SH2 DOMAIN	ENT COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	ENT COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	FERASE NT UNIT ICMK MK 4	HOSPHOTR MP- NTEIN
Coumpound	TYPE I; CHAIN: B, D, F, H;	FKS06-BINDING PROTEIN; CHAIN: A, C, E, G; TGF-B SUPERFAMILY RECEPTOR TYPE I; CHAIN: B, D, F, H;	PP60 V-SRC TYROSINE KINASE TRANSFORMING PROTEIN, CHAIN: NULL;	PP60 V-SRC TYROSINE KINASE TRANSFORMING PROTEIN; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37) ICMK 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN
SeqFold score				165.48		99.54		99.49
PMF		1	_					
Verify score		0.51	1.13		99.0		0.7	
PSI- BLAST		2.60E-74	2.60E-40	2.60E-40	1.20E-34	1.20E-34	9.10E-35	6.50E-35
End		531	259	263	529	538	515	542
Start		256	155	155	269	270	282	227
Chain ID		В			<	<	ப	<u>ш</u>
PDB ID		1b6c	1 bk l	1 bk	1blx	1blx	Lcm k	1ctp
SEQ ID		607	209	209	607	607	209	209

PDB annotation				TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER		PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE	PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS PHOSPHORYI ATION	PROTEIN KINASE CDK2;	TRANSFERASE, SERINE/THREONINE	PROTEIN KINASE, ATP-BINDING, 2	MITOSIS, PHOSPHORYLATION	TRANSFERASE INK3; TRANSFERASE,	JONES MAP KINASE, SERINE/THREONINE PROTEIN 2 KINASE	COMPLEX (TRANSFERASE/PEPTIDE) SRC, SH3 DOMAIN, LIGANDS, NON-	PEPTIDE ELEMENTS, Z COMPLEX (TRANSFERASE/PEPTIDE)	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE:	TRANSFERASE, MAP KINASE, SERINE/THREONINE-PROTEIN	KINASE, 2 P38			
Coumpound	SUBUNIT) ICTP 4	TRANSFERASE(PHOSPHOTR ANSFERASE) CAMP- DEPENDENT PROTEIN	KINASE (E.C.Z.7.1.37) (CAPR) ICTP 3 (CATALYTIC SUBUNIT) ICTP 4	SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B;	SERINE/THREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: C, D;	HUMAN CYCLIN- DEPENDENT KINASE 2;	CHAIN: NULL;	HUMAN CYCLIN-	DEPENDENT KINASE 2;	CHAIN: NULL;		C-JUN N-TERMINAL	KINASE; CHAIN: NULL;	C-SRC; CHAIN: C; NLI (MN7- MN2-MN1-PLPPLP); CHAIN:	. .	MAP KINASE P38; CHAIN:			PHOSPHOTRANSFERASE V- SRC TYROSINE KINASE	TRANSFORMING PROTEIN (PHOSPHOTYROSINE 1SHA 3	RECOGNITION DOMAIN
SeqFold score						111.67													165.79		
PMF		_		_				-				1		-		-					
Verify score		0.55		0.71				0.43				9.0		0.13		0.44					
PSI- BLAST		6.50E-35		1.30E-42		1.30E-37		1.30E-37				1.30E-33		2.60E-19		2.60E-32			9.10E-40		
End		515		519		538		529				538		149		517			256		
Start		282		272		273		281				269		88		269			154		
Chain ID		ш		၁										ی					٧		
PDB ID		lctp		1f3m		Ihcl		The				ljnk		Inlo		1p38			1sha		
SEQ NO.		209		209		607		209				209		209		209			209		

# K	Start End AA AA	H PSI-	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
						SH2) (E.C.2.7.1.112) COMPLEX WITH 1SHA 4 PHOSPHOPEPTIDE A (TYR- VAL-PRO-MET-LEU, PHOSPHORYLATED TYR) 1SHA 5	
154	256	9.10E-40	0.97	_		PHOSPHOTRANSFERASE V-SRC TYROSINE KINASE TRANSFORMING PROTEIN (PHOSPHOTYROSINE ISHA 3) (PHOSPHOTYROSINE ISHA 3) (PECCGNITION DOMAIN SH2) (E.C.2.7.1.112) COMPLEX WITH ISHA 4 PHOSPHOPEPTIDE A (TYR-VAL-PRO-MET-LEU, PHOSPHORYLATED TYR) ISHA 5	
100	166	1.30E-14	-0.02	0.89		ALPHA-SPECTRIN; CHAIN: NULL;	CYTOSKELETON CAPPING PROTEIN, CALCIUM-BINDING, DUPLICATION, REPEAT, 2 SH3 DOMAIN, CYTOSKELETON
	130	3.90E-09	-0.25	0.01		ALPHA-SPECTRIN; CHAIN: NULL;	CYTOSKELETON CAPPING PROTEIN, CALCIUM-BINDING, DUPLICATION, REPEAT, 2 SH3 DOMAIN, CYTOSKELETON
	255	9.10E-34			121.36	ABL TYROSINE KINASE; CHAIN: NULL;	TRANSFERASE TRANSFERASE, TYROSINE KINASE, SH3, SH2, ONCOPROTEIN
	255	9.10E-34	29.0	_		ABL TÝROSINE KINASE; CHAIN: NULL;	TRANSFERASE TRANSFERASE, TYROSINE KINASE, SH3, SH2, ONCOPROTEIN
	425	6.50E-59	-0.13	0.54		SHP-2; CHAIN: A, B;	TYROSINE PHOSPHATASE SYP, SHPTP-2; TYROSINE PHOSPHATASE, INSULIN SIGNALING, SH2 PROTEIN
203	284	9.106-21	-0.13	0.31		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
509	280	1.30E-32	0	0.74		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC

PDB annotation	FINGER, DNA-BINDING PROTEIN			COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDNA) ZINC FINGER. PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Coumpound	OLIGONUCLEOTIDE BINDING SITE, CHAIN. B, C,	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR ADRI (RESIDUES 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADRIB) 1ARD 5	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SeqFold								
PMF score		0.05	0.17	0.87	-	_	-	_
Verify score		-0.34	-0.3	0.16	0.32	0.89	0.67	0.62
PSI- BLAST		6.50E-05	2.60E-16	1.30E-22	6.50E-33	1.20E-42	5.20E-44	9.10E-47
End		537	537	283	311	339	367	395
Start		209	485	202	230	259	286	314
Chain ID				O	၁	ပ	၁	U
PDB ID		lard	1990	I me	Ime y	lmc y	Ime y	lme y
SEQ	Ö	809	809	809	809	809	809	809

PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
	<u>e</u>		V	BLASI	score	score	score		
									(ZINC FINGER/DNA)
E	O	314	396	9.10E-47			102.94	DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER PROTEIN-DNA
	- <u></u>							PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
									CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
	C	342	423	7 80F-46	0.74	_		DNA: CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
<u>}</u>		!	!					CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
								PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
									CRYSIAL SIRUCIONE, COMPLEX (ZINC FINGENDNA)
4	O	370	451	1.30E-45	0.53	-		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
,								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
								PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
									CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
ì e	ر	308	479	1 30F-40	0 33	_		DNA CHAIN A. B. D. E.	COMPLEX (ZINC FINGER/DNA) ZINC
ر : ک		2	<u>`</u>		3	•		CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
								PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
									CRYSTAL STRUCTURE, COMPLEX
- 1									(ZINC FINGER/DNA)
Jne E	ပ	426	202	2.60E-34	0.05			DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGERODNA) ZINC
								CONSENSUS ZINC FINGER	FINGER, FROI EJINEDIA INTERACTION PROTEIN DESIGN 2
								PROTEIN; CHAIN: C, F, G;	CRYSTAL STRICTURE COMPLEX
									(ZINC FINGER/DNA)
E	O	454	589	3.90E-32	-0.08	0.22		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
								PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
									CRISIAL SINOCIOIG, COMPLEX (ZINC FINGER/DNA)
1 2	J	536	617	3.90E-42	0.2	_		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC
>								CONSENSUS ZINC FINGER	FINGER, PROTEIN-DNA
								PROTEIN; CHAIN: C, F, G;	INTERACTION, PROTEIN DESIGN, 2
		<u>.</u> -							CRYSTAL STRUCTURE, COMPLEX
ŀ	+	;				\ -\		4	COM THE CARLO ENICED (DNA) ZINC
Ĕ.	ပ	564	645	3.90E-46	0.29			DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGENDINA) ZINC
								PROTEIN CHAIN C. F. G.	INTERACTION, PROTEIN DESIGN, 2
	_								CRYSTAL STRUCTURE, COMPLEX
- 1									(ZINC FINGER/DNA)

PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ZINC FINGER TRANSCRIPTION FACTOR SPI; ZINC FINGER, TRANSCRIPTION ACTIVATION, SPI	ZINC FINGER TRANSCRIPTION FACTOR SPI; ZINC FINGER, TRANSCRIPTION ACTIVATION, SPI	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION NITIATION, ZINC FINGER PROTEIN		COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN
	CRY CRY	CON FINC INTE CRY	FAC	ZING FAC TRA	CON REG TRA		TRA TRA FINC FINC TRA	N TEG
Coumpound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	SP1F3; CHAIN: NULL;	SPIF2; CHAIN: NULL;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TRANSCRIPTION REGULATION TRANSCRIPTIONAL ELONGATION FACTOR SII (TFIIS, NUCLEIC-ACID 1TFI 3 BINDING DOMAIN) (NMR, 12 STRUCTURES) 1TFI 4	YYI; CHAIN: C. ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold					109.61			
PMF	1	-	0.01	0.13		0.24	0.03	-
Verify	0.33	9.0	-0.29	-0.23		-0.22	-0.34	0.26
PSI. BLAST	2.60E-46	2.60E-45	7.80E-05	9.10E-05	2.60E-71	0.0091	7.80E-29	6.50E-43
End	673	702	537	537	505	520	311	340
Start	592	620	509	\$00	342	480	165	229
Chain ID	O	O			∢		ပ	ပ
PDB ID	Jme y	1me y	lgl	1sp2	1116	91	lubd	pqn1
SEQ ID NO:	809	809	809	809	809	809	809	809

1D 1D AA BLAST Store	ξĞ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
Ubd C 263 368 780E-32 0.55 1 YY1; CHAIN; C; ADENO- C C C C C C C C C	<u> </u>	<u> </u>	E	VV	VV V	BLAST	score	score	score		
1ubd C 263 368 780E-32 0.55 1 YYI; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: C. ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO-ASSOCIATED VIRIUS PS INITIATOR ELEMENT DNA; CHAIN: A, B											RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Lubd C 284 395 1.30E-53 0.58 1 TYYI; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: G, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN:	809	1ubd	ပ	263	368	7.80E-52	0.55	_		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA:	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,
Lubd C 284 395 1.30E-53 0.58 1 TVYI; CHAIN: C; ADENO- CHAIN: A, B; CHAIN: C; ADENO- CHAIN: C; ADENO- CHAIN: C, ADENO- C, C, C, C, C, C, C, C, C, C, C, C, C,										CHAIN: A, B;	INITIATOR ELEMENT, YYI, ZINC 2
Lubd C 284 395 1.30E-53 0.58 1 TYY1; CHAIN; C; ADENO- ASSOCIATED VIRUS PS											RECOGNITION, 3 COMPLEX
Lubd C 341 451 5.20E-53 0.26 1 YYY1; CHAIN: C. ADENO- CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C. ADENO- CHAIN: A, B; CHAIN: C. ADENO- CHAIN: A, B; C. CHAIN: C. ADENO- CHAIN: A, B; C. CHAIN: A, B; C. CHAIN: A, B; C. CHAIN: C. ADENO- C. C. C. C. C. C. C. C. C. C. C. C. C.	8	Lubd	C	284	395	1 30E-53	0.58	-		YY1: CHAIN: C: ADENO-	COMPLEX (TRANSCRIPTION
Lubd C 341 451 5.20E-53 0.26 1 YY1; CHAIN; C; ADENO- ASSOCIATED VIRUS P5 Initiator element DNA; CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- CHAIN C, ADENO- C, C, C, C, C, C, C, C, C, C, C, C, C,)		}			•		ASSOCIATED VIRUS P5	REGULATION/DNA) YING-YANG 1;
lubd C 341 451 5.20E-53 0.26 1 YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS P5	-									CHAIN: A, B;	INTIATOR ELEMENT, YYI, ZINC 2
Lubd C 341 451 5.20E-53 0.26 1 ASSOCIATED VIRUS P5											FINGER PROTEIN, DNA-PROTEIN
1ubd C 341 451 5.20E-53 0.26 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 374 479 9.10E-51 0.34 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 424 589 2.60E-39 -0.28 0.34 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 XY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 ASSOCIATED VIRUS P5 Iuitiator Element DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Iuitiator Element DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Iuitiator Element DNA; CHAIN: C, ADENO-CHAIN:											RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
1ubd C 374 479 9, 10E-51 0,34 1 YY1; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iubd C 424 589 2,60E-39 -0,28 0,34 YY1; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iubd C 424 589 2,60E-39 -0,28 0,34 YY1; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iubd C 508 618 5,20E-45 -0,04 0,99 YY1; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iuritator element DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Initiator element DNA; CHAIN: A, B; CHAI	809	lubd	၁	341	451	5.20E-53	0.26	-		YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION DECIT A TICKNON A) VING VANG I
1ubd C 374 479 9.10E-51 0.34 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Initiator ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Initiator ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Initiator ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Initiator ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Initiator ELEMENT DNA; CHAIN: A, B; CHA										NITIATOR ELEMENT DNA;	TRANSCRIPTION INITIATION,
1ubd C 374 479 9.10E-51 0.34 1 TYY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiator element display CHAIN: A, B; CHAIN: A, B; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 TYY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 TYY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiator element display CHAIN: A, B; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiator element display CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: C; ADENO-CH		_				_				CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2
1ubd C 374 479 9.10E-51 0.34 1 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 1ubd C 424 589 2.60E-39 -0.28 0.34 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5											FINGER PROTEIN, DNA-PROTEIN
1ubd C 374 479 9.10E-51 0.34 1 YY1; CHAIN: C. ADENO-ASSOCIATED VIRUS P5 1ubd C 424 589 2.60E-39 -0.28 0.34 YY1; CHAIN: C. ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C. ADENO-ASSOCIATED VIRUS P5 1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C. ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C. ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C. ADENO-ASSOCIATED VIRUS P5 Intransport CHAIN: A. B. CHAIN: A. B. CHAIN: A. B. CHAIN: A. B.											RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGISTATION/DNA)
1ubq C 424 589 2.60E-39 -0.28 0.34 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 NITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 NITIATOR ELEMENT DNA; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 NITIATOR ELEMENT DNA; CHAIN: C, ADENO-CHAIN: A, B; CHAIN: C, ADENO-CHAIN: A, B; CHAIN: C, ADENO-CHAIN: A, B; CHAIN: C, ADENO-CHAIN: C, ADENO-CHAIN: A, B; CHAIN: C, ADENO-CHAIN: C, ADENO-CHAIN: C, ADENO-CHAIN: C, ADENO-CHAIN: A, B; CHAIN: C, ADENO-CHAIN: C,	80	lubd	ပ	374	479	9.10E-51	0.34	_		YY1: CHAIN: C. ADENO-	COMPLEX (TRANSCRIPTION
Iubd C 424 589 2.60E-39 -0.28 0.34 YYI; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YYI; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iuriator Element DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Initiator Element DNA; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Initiator Element DNA; CHAIN: C, ADENO- CHAIN: A, B; Initiator Element DNA; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- CHAIN: C, ADENO- CHAIN: A, B; CHAIN: C, ADENO- C, CHAIN: C, ADENO- C, CHAIN: C,								•		ASSÓCIATED VIRUS PS	REGULATION/DNA) YING-YANG 1;
Iubd C 424 589 2.60E-39 -0.28 0.34 YYI; CHAÏN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YYI; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Initiator Element DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B;										INITIATOR ELEMENT DNA;	TRANSCRIPTION INITIATION,
Iubd C 424 589 2.60E-39 -0.28 0.34 YY1; CHAÏN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5 Inbd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C, ADENO-ASSOCIATED VIRUS P5										CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2
lubd C 424 589 2.60E-39 -0.28 0.34 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Inbd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Inbd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Inititation Element DNA; CHAIN: A, B: CHAIN: A, B: CHAIN: A, B: CHAIN: A, B:											FINGER PROTEIN, DNA-PROTEIN RECOGNITION 3 COMPLEX
lubd C 424 589 2.60E-39 -0.28 0.34 YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiation Element DNA; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiation Element DNA; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B;											(TRANSCRIPTION REGULATION/DNA)
NITIATOR ELEMENT DNA; CHAIN: A, B; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 Iubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C, ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B.	808	lubd	၁	424	589	2.60E-39	-0.28	0.34		YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION DECLIT ATTIONIAN VINC MANG 1:
CHAIN: A, B; C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B.										NITIATOR ELEMENT DNA;	TRANSCRIPTION INITIATION,
lubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiation Element DNA; Initiation Element DNA; CHAIN: A. B.										CHAIN: A, B;	INITIATOR ELEMENT, YY1, ZINC 2
1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 Initiation Element DNA; Initiation Element DNA; CHAIN: A. B.											FINGER PROTEIN, DNA-PROTEIN
1ubd C 508 618 5.20E-45 -0.04 0.99 YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A. B.											KECOGNITION, 3 COMPLEX (TRANSCRIPTION REGIT ATTON/DNA)
	80	lubd	၁	808	819	5.20E-45	-0.04	0.99		YY1; CHAIN: C; ADENO-	COMPLEX (TRANSCRIPTION
					الدائد المالية					ASSOCIATED VIRUS P5	REGULATION/DNA) YING-YANG I,
										CHAIN A B.	IRANSCRIPTION INITIATION, INITIATOR ELEMENT VVI 21NC 2

PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADR1, ZINC FINGER, NMR			COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	╀┤
Coumpound		YYI; CHAIN: C: ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A. B;	YYI; CHAIN: C, ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	ADRI; CHAIN: NULL;	COMPLEX(TRANSCRIPTION REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX(TRANSCRIPTION REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII;
SeqFold score									
PMF		-		0.41	90.0	0.23	-	-	
Verify score		-0.07	0.06	-0.15	0.11	0.28	0.1	0.45	0.32
PSI- BLAST		5.20E-56	1.30E-54	2.60E-18	1.30E-15	6.50E-18	9.10E-58	1.00E-65	1.30E-67
End AA		674	102	509	530	563	369	397	480
Start		564	590	455	479	505	230	259	314
Chain ID		O	O		<	<	<	A	4
PDB		lubd	pqnl	2adr	2drp	2drp	2gli	2gli	2gli
SEQ 15		809	809	809	809	809	809	809	809

PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
								CHAIN: A, DNA; CHAIN: C, D;	PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A		398	619	1.30E-56	0	0.37		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)
2gli A		536	675	1.30E-64	0.3	_		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA: CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A		564	701	2.60E-70	0.16	0.99		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A		564	703	2.60E-70			92.71	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
lam D		9	165	2.60E-20	0.46	-		P50-RHOGAP; CHAIN: A, B, C; CDC42HS; CHAIN: D, E, F;	COMPLEX (GTPASE-ACTIVATING/GTP-BINDING) COMPLEX (GTPASE-ACTIVATING/GTP-BINDING), GTPASE ACTIVATION
lbyu B			192	3.90E-24			55.55	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	TRANSPORT PROTEIN TC4; GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN
lbyu B		9	181	3.90E-24	0.15	0.66		GTP-BINDING PROTEIN RAN: CHAIN: A, B;	TRANSPORT PROTEIN TC4, GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN
lcxz A			187	1.30E-20			61.64	HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B;	SIGNALING PROTEIN PROTEIN- PROTEIN COMPLEX, ANTIPARALLEL COILED-COIL
lcxz A		9	0.71	1.30E-20	0.19	-		HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B;	SIGNALING PROTEIN PROTEIN- PROTEIN COMPLEX, ANTIPARALLEL COILED-COIL
1 ds6 A		9	172	2.60E-20	0.14	_		RAS-RELATED C3 BOTULINUM TOXIN	SIGNALING PROTEIN P21-RAC2; RHO GDI 2, RHO-GDI BETA, LY-GDI; BETA

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
ΒŞ	9	e	V V	¥	BLAST	score	score	score		
Ċ									SUBSTRATE 2; CHAIN: A; RHO GDP-DISSOCIATION	SANDWHICH, PROTEIN-PROTEIN COMPLEX, G-DOMAIN, 2
									INHIBITOR 2; CHAIN: B;	IMMUNOGLOBULIN FOLD, WALKER FOLD, GTP-BINDING PROTEIN
609	1e0s	A	9	891	1.20E-22	0.12	0.57		ADP-RIBOSYLATION	G PROTEIN G PROTEIN, RAS, ARF,
									FACTOR 6; CHAIN: A;	ARF6, MEMBRANE TRAFFIC
609	Ihur	٧	_	172	7.80E-23			55.9	HUMAN ADP-	PROTEIN TRANSPORT GDP-BINDING,
									RIBOSYLATION FACTOR 1;	MEMBRANE TRAFFICKIN, NON-
									1HUR 5 CHAIN: A, B; 1HUR 7	MYRISTOYLATED IHUR 16
609	Ihur	¥	9	168	7.80E-23	-0.12	0.55		HUMAN ADP-	PROTEIN TRANSPORT GDP-BINDING.
									RIBOSYLATION FACTOR 1;	MEMBRANE TRAFFICKIN, NON- MYRISTOYI ATED IHUR 16
609	l.hr	4	3	174	3.90E-23			71.05	RAN: CHAIN: A. C.	SMALL GTPASE KARYOPHERIN
-	=	;	`	· · · ·					IMPORTIN BETA SUBUNIT:	BETA, P95 SMALL GTPASE, NUCLEAR
									CHAIN: B, D;	TRANSPORT RECEPTOR
609	1 ibr	٧	9	173	3.90E-23	0.04	0.99		RAN; CHAIN: A, C;	SMALL GTPASE KARYOPHERIN
									IMPORTIN BETA SUBUNIT;	BETA, P95 SMALL GTPASE, NUCLEAR
									CHAIN: B, D;	TRANSPORT RECEPTOR
609	lmh		2	681	3.90E-21			52.65	RACI; CHAIN: NULL;	GTP-BINDING GTP-BINDING, GTPASE,
	_									SMALL G-PROTEIN, KHO FAMILY,
			,];		ì			THE LEGISLE	CTE PRIDATE CTE PRIDAIC CTPACE
609	Ē.		9	=======================================	3.90E-21	0.36	-		KACI; CHAIN: NULL;	CIF-BINDING CIF-BINDING, CIT ASE, CMAIL G-PROTEIN RHO FAMILY
	_		-							RAS SUPER 2 FAMILY
90	1	ļ	,	107	2010			97.77	DANI CHAMI: A C. MITCI EAD	COMPLEX (SMALL GTPASE/NITCLEAR
600	di I	ر	<u>-</u>	\ <u>0</u>	9.10E-24			0/://	PORE COMPLEX PROTEIN	PROTEIN) COMPLEX (SMALL
									NTP358: CHAIN: B. D:	GTPASE/NUCLEAR PROTEIN), SMALL
		_								GTPASE, 2 NUCLEAR TRANSPORT
609	lπp	U	9	180	9.10E-24	-0.02	9.0		RAN; CHAIN: A, C; NUCLEAR	COMPLEX (SMALL GTPASE/NUCLEAR
									PORE COMPLEX PROTEIN	PROTEIN) COMPLEX (SMALL
									NUP358; CHAIN: B, D;	GIPASE/NUCLEAK PROTEIN), SMALL
8	1	-	,	301	00 307 0			36 83	DSO BUOGAB: CUAIN: A	COMPLEX GTPASE
600	*×1	n 	·	6	7.00E-20			74.33	TOUR ANSENDAMENT DE ANSENDAMENT	ACTIVATIVEROTO-ONCOGENE)
									RHOA: CHAIN: B:	GTPASE-ACTIVATING PROTEIN
_										RHOGAP; COMPLEX (GTPASE
										ACTIVATION/PROTO-ONCOGENE),
										GTPASE, 2 TRANSITION STATE, GAP
609	1tx4	В	5	165	2.60E-20	0.37	-		P50-RHOGAP, CHAIN: A;	COMPLEX(GTPASE
									TRANSFORMING PROTEIN	ACTIVATN/PROTO-ONCOGENE)
									RHOA; CHAIN: B;	GIPASE-ACTIVATING PROTEIN

PDB	8 -	Chain	Start	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
										RHOGAP; COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP
ylq1		<	370	169	2.60E-52			129.83	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA. COMPLEX (INHIBITOR PROTEIN/KINASE)
1hcl	-		371	687	7.80E-50			118.75	HUMAN CYCLIN. DEPENDENT KINASE 2; CHAIN: NULL;	PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP-BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION
layz	2.	∢	167	337	1.00E-15			54.69	UBIQUITIN-CONJUGATING ENZYME RAD6; CHAIN: A, B, C;	UBIQUITIN CONJUGATION UBC2; UBIQUITIN CONJUGATION, UBIQUITIN-CONJUGATING ENZYME
1959	8	¥	168	335	2.60E-17			54.87	UBIQUITIN CONJUGATING ENZYME; CHAIN: A;	LIGASE UBIQUITIN, UBIQUITIN- CONJUGATING ENZYME, YEAST
2aak	¥		191	333	2.60E-14			59.33	UBIQUITIN CONJUGATING ENZYME; CHAIN: NULL;	UBIQUITIN CONJUGATION UBCI; UBIQUITIN CONJUGATION, LIGASE
laj4	4		317	468	5.20E-06			70.38	TROPONIN C, CHAIN: NULL;	MUSCLE PROTEIN CTNC; CARDIAC, MUSCLE PROTEIN, REGULATORY, CALCIUM BINDING
laui	-5	<u>a</u>	320	476	7.80E-21			78.45	SERINE/THREONINE PHOSPHATASE 2B; CHAIN: A, B;	HYDROLASE CALCINEURIN; HYDROLASE, PHOSPHATASE, IMMUNOSUPPRESSION
3	1 ika		311	491	5.20E-20			67.12	RECOVERIN; CHAIN: NULL;	CALCIUM-BINDING PROTEIN CALCIUM-MYRISTOYL SWITCH, CALCUIM-BINDING PROTEIN
=	Itef		320	465	9.10E-17			68.38	TROPONIN C; CHAIN: NULL;	CALCIUM-REGULATED MUSCLE CONTRACTION MUSCLE CONTRACTION, CALCIUM-BINDING, TROPONIN, E-F HAND, 2 OPEN CONFORMATION REGULATORY DOMAIN, CALCIUM-REGULATED 3
- Itx	ž		320	463	1.20E-16			67.26	TROPONIN C; 1TNX 4	CALCIUM-BINDING PROTEIN EF-

	П			10	ISE	O	9 HO	GTP	EL	HO JR
PDB annotation	HAND ITNX 14	PHOSPHOTRANSFERASE PHOSPHOTRANSFERASE		PHOSPHOTRANSFERASE NUCLEOSIDE TRIPHOSPHATE, NUCLEOSIDE DIPHOSPHATE INUE 10	COMPLEX (GTPASE- ACTIVATING/GTP-BINDING) COMPLEX (GTPASE- ACTIVATING/GTP-BINDING), GTPASE ACTIVATION	SIGNALING PROTEIN GTP-BINDING PROTEINS, PROTEIN-PROTEIN COMPLEX, EFFECTORS	SIGNALING PROTEIN GTP-BINDING PROTEIN RHOA, GTPASE RHOA; RHO GDI 1; RHO GTPASE, G-PROTEIN, SIGNALING PROTEIN	SIGNALING PROTEIN G PROTEIN. GTP HYDROLYSIS, KINETIC CRYSTALLOGRAPHY, 2 SIGNALING PROTEIN	SIGNALING PROTEIN PROTEIN- PROTEIN COMPLEX, ANTIPARALLEL COILED-COIL	SIGNALING PROTEIN P21-RAC2; RHO GDI 2, RHO-GDI BETA, LY-GDI; BETA SANDWHICH, PROTEIN-PROTEIN COMPLEX, G-DOMAIN, 2 IMMUNOGLOBULIN FOLD, WALKER FOLD, GTP-BINDING PROTEIN
	HAN	PHOS		PHOS NUCI NUCI	COMI COMI ACTI	SIGN PROT COMI	PROT GDI I SIGN	SIGNALIN HYDROL CRYSTAL PROTEIN	SIGN PROT COIL	SIGN GDI 2 SAND COMI IMMI
Coumpound	CHAIN: NULL; 1TNX 5	NUCLEOSIDE DIPHOSPHATE TRANSFERASE; CHAIN: A, B, C;	PHOSPHOTRANSFERASE NUCLEOSIDE DIPHOSPHATE KINASE (E.C.2.7.4.6) INSQ 3	NUCLEOSIDE DIPHOSPHATE KINASE; INUE 4 CHAIN: A, B, C, D, E, F; INUE 5	P50-RHOGAP; CHAIN: A, B, C; CDC42HS; CHAIN: D, E, F;	RAS-RELATED PROTEIN RAP-1A; CHAIN: A; PROTO- ONKOGENE SERINE/THREONINE PROTEIN KINASE CHAIN: B;	TRANSFORMING PROTEIN RHOA; CHAIN: A, C, RHO GDP DISSOCIATION INHIBITOR ALPHA; CHAIN: E, F;	TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A;	HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181), CHAIN: A; PKN; CHAIN: B;	RAS-RELATED C3 BOTULNUM TOXIN SUBSTRATE 2; CHAIN: A; RHO GDP-DISSOCIATION INHIBITOR 2; CHAIN: B;
SeqFold score		105.78	104.27	105.45						
PMF					0.55	0	0.65	0.36	0.93	0.54
Verify score					0.31	0.17	0.24	0.23	0.39	0.05
PSI- BLAST		3.60E-53	3.60E-50	9.00E-53	3.60E-56	5.40E-56	9.00E-62	1.80E-59	9.00E-62	1.40E-62
End		115	115	116	263	264	271	265	264	267
Start AA			_	_	88	98	68	98	68	88
Chain ID		¥.	A	V	D	¥	V	V	A	¥
PDB ID		lbe4	lnsq	lnue	lam 4	lcly	1cc0	lctq	zxɔj	1ds6
SEQ ID NO:		646	646	646	859	859	859	859	859	658

(<u>)</u>	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
} ∣		9	Ş	AA	DLASI	score	score	3008		
1 hur		V	75	266	7.20E-11			54.59	HUMAN ADP- RIBOSYLATION FACTOR 1; IHUR 5 CHAIN: A, B; IHUR 7	PROTEIN TRANSPORT GDP-BINDING, MEMBRANE TRAFFICKIN, NON- MYRISTOYLATED IHUR 16
ᄪ	£		88	267	1.30E-63	0.19	0.45		RACI; CHAIN: NULL;	GTP-BINDING GTP-BINDING, GTPASE, SMALL G-PROTEIN, RHO FAMILY, RAS SUPER 2 FAMILY
154	4	B	68	263	1.80E-58	0.31	0.82		PSO-RHOGAP, CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;	COMPLEX(GTPASE ACTIVATIVPROTO-ONCOGENE) GTPASE-ACTIVATING PROTEIN RHOGAP; COMPLEX (GTPASE ACTIVATION/PROTO-ONCOGENE), GTPASE 2 TRANSITION STATE GAP
2ngr		V	88	270	7.20E-60	-0.01	0.4		GTP BINDING PROTEIN (025K); CHAIN: A; GTPASE ACTIVATING PROTEIN (RHG); CHAIN: B;	HYDROLASE CDC42/CDC42GAP: CDC42/CDC42GAP; TRANSITION STATE, G-PROTEIN, GAP, CDC42, ALF3., HYDROLASE
i 🕾 🔝 🖠	Згар	Y	98	265	1.10E-54	0.11	0.47		RAB3A; CHAIN: A;	HYDROLASE G PROTEIN, VESICULAR TRAFFICKING, GTP HYDROLYSIS, RAB 2 PROTEIN, NEUROTRANSMITTER RELEASE, HYDROLASE
آھ: ا	lbor		91	59	9.00E-06	-0.73	0.25		TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION
ਹ ।	1chc		16	09	9.00E-15	0.24	86:0		VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	
[€	1fbv	V	<u>&</u>	99	1.805-10	0.25	6.0		SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
(60)	1g25	∢	8-	72	1.30E-05	-0.44	60.0		CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)
	1pjr		172	218	0.002	-0.75	0.21		PCRA; CHAIN: NULL;	HELICASE DNA REPAIR, DNA

PDB annotation	REPLICATION, SOS RESPONSE, HELICASE, 2 ATP-BINDING, DNA- BINDING	HELICASE DNA REPAIR, DNA REPLICATION, SOS RESPONSE, HELICASE, ATP- 2 BINDING, DNA- BINDING	COMPLEX (HELICASE/DNA) COMPLEX (HELICASE/DNA), HELICASE, DNA UNWINDING	HYDROLASEDNA ATP-DEPENDENT HELICASE PCRA; ATP-DEPENDENT HELICASE PCRA; HELICASE PCRA, HYDROLASE, DNA, PRODUCT COMPLEX	TRANSCRIPTION HELIX-BUNDLE	TRANSCRIPTION HELIX-BUNDLE			
Coumpound		PCRA (SUBUNIT); CHAIN: A; PCRA (SUBUNIT); CHAIN: B; PCRA (SUBUNIT); CHAIN: C; PCRA (SUBUNIT); CHAIN: D;	ATP-DEPENDENT DNA HELICASE REP; CHAIN: A, B; DNA CHAIN: C;	HELICASE PCRA; CHAIN: A, F; HELICASE PCRA; CHAIN: B, G; DNA (5'-D(*TP*TP*TP*T)-3'); CHAIN: C, D; DNA (5'-D(*GP*C)-3'); CHAIN: H; DNA (5'-D(*AP*CP*TP*GP*C)-3'); CHAIN: I;	TRANSCRIPTION ELONGATION FACTOR S-II; CHAIN: A;	TRANSCRIPTION ELONGATION FACTOR S-II; CHAIN: A;	TRANSCRIPTION REGULATION TRANSCRIPTIONAL ELONGATION FACTOR SII (TFIIS, NUCLEIC-ACID 1TFI 3 BINDING DOMAIN) (NMR, 12 STRUCTURES) 1TFI 4	TRANSCRIPTION REGULATION TRANSCRIPTIONAL ELONGATION FACTOR SII (TFIIS, NUCLEIC-ACID 1TFI 3 BINDING DOMAIN) (NMR, 12 STRUCTURES) 1TFI 4	TRANSCRIPTION REGULATION
SeqFold score							:	84.88	
PMF		0.28	69:0	0.88	0.24	0.18	-		
Verify score		-0.84	-0.67	-0.31	-0.43	0.45	0.45		0.45
PSI- BLAST		0.0045	0.00011	9.00E-06	1.00E-18	3.60E-12	1.30E-22	3.90E-24	3.90E-24
End AA		202	238	236	49	64	332	332	332
Start		172	172	172	_	4	283	283	283
Chain ID		В	4	∢	V	¥			
PDB ID		1qhg	luna	2pjr	1 c 00	1co0	1tfi	Jil I	144
SEQ FD		661	199	199	663	663	663	663	663

1 1	Start AA	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
							ELONGATION FACTOR SII (TFIIS, NUCLEIC-ACID ITFI 3 BINDING DOMAIN) (NMR, 12 STRUCTURES) ITFI 4	
388		452	1.60E-13	90.0	0.99		DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B; RNA (5'- R(*GP*GP*CP*GP*CP *GP*CP*C)-3'); CHAIN: C. D, E, G;	RNA BINDING PROTEINRNA XLRBPA, PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA
390		451	1.30E-14	0.45	0.42		DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B; RNA (5'- R(*GP*GP*CP*GP*CP *GP*CP*C)-3'); CHAIN: C, D, E, G;	RNA BINDING PROTEIN/RNA XLRBPA, PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA
512		574	2.60E-16	0.11	66.0		DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B; RNA (5'- R(*GP*GP*CP*GP*CP *GP*CP*C)-3'); CHAIN: C, D, E, G;	RNA BINDING PROTEIN/RNA XLRBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA
514		558	1.80E-09	-0.16	0.39		DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B; RNA (5'- R(*GP*GP*CP*GP*CP*GP*CP *GP*CP*C)-3'); CHAIN: C, D, E, G;	RNA BINDING PROTEIN/RNA XLRBPA, PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA
388		452	1.60E-07	90.0	0.7		DOUBLE STRANDED RNA BINDING PROTEIN A; CHAIN: A, B; RNA (5'- R(*GP*GP*CP*GP*CP*GP*CP *GP*CP*C)-3'); CHAIN: C, D, E, G;	RNA BINDING PROTEIN/RNA XLRBPA; PROTEIN-RNA COMPLEX, DOUBLE STRANDED RNA, PROTEIN- RNA 2 INTERACTIONS, RNA-BINING PROTEIN, RNA BINDING PROTEIN/RNA
380		454	1.80E-14	-0.12	0.31		MATERNAL EFFECT PROTEIN (STAUFEN); CHAIN: A: STAUFEN DOUBLE-STRANDED RNA BINDING DOMAIN; CHAIN:	CELL CYCLERNA DSRBDIII; NMR STRUCTURE, PROTEIN/RNA, PROTEIN DSRBD. DROSOPHILA. RNA 2 HAIRPIN

Verify PMF SeqFold Coumpound PDB annotation score score score	-0.06 0.69 MATERNAL EFFECT CELL CYCLERNA DSRBDIII; NMR PROTEIN (STAUFEN); STRUCTURE, PROTEIN/RNA, PROTEIN CHAIN: A; STAUFEN DSRBD, DROSOPHILA, RNA 2 DOUBLE-STRANDED RNA HAIRPIN BINDING DOMAIN; CHAIN: B:	0.04 0.39 MATERNAL EFFECT CELL CYCLERNA DSRBDIII; NMR PROTEIN (STAUFEN); STRUCTURE, PROTEIN/RNA, PROTEIN CHAIN: A; STAUFEN DSRBD, DROSOPHILA, RNA 2 DOUBLE-STRANDED RNA HAIRPIN BINDING DOMAIN; CHAIN: B;	0.75 1 MATERNAL EFFECT CELL CYCLE/RNA DSRBDIII; NMR PROTEIN (STAUFEN); STRUCTURE, PROTEIN/RNA, PROTEIN CHAIN: A; STAUFEN DSRBD, DROSOPHILA, RNA 2 DOUBLE-STRANDED RNA HAIRPIN BINDING DOMAIN; CHAIN: B:	-0.17 0.06 PROTEIN KINASE PKR; TRANSFERASE DSRNA-BINDING CHAIN: A; DOMAIN, NMR, PKR, SOLUTION STRUCTURE, PROTEIN 2 KINASE. TRANSFERASE	-0.05 0.09 PROTEIN KINASE PKR; TRANSFERASE DSRNA-BINDING CHAIN: A: DOMAIN, NMR, PKR, SOLUTION STRUCTURE, PROTEIN 2 KINASE, TRANSFERASE	0.23 0.07 MATERNAL EFFECT DOUBLE STRANDED RNA BINDING PROTEIN STAUFEN; 1STU 4 DOMAIN STAUFEN 1STU 13	0.19 1 MATERNAL EFFECT DOUBLE STRANDED RNA BINDING PROTEIN STAUFEN; ISTU 4 DOMAIN STAUFEN ISTU 13		0.48 0.92 SXL-LETHAL PROTEIN; RNA-BINDING PROTEIN/RNA TRA CHAIN: A, B; RNA (5'- PRE-MRNA; SPLICING REGULATION, RP-GP*UP*UP*UP*UP*UP*UP*U RNP DOMAIN, RNA COMPLEX
PSI- BLAST	5.20E-15	5.40E-05	1.30E-19	1.10E-15	7.20E-08	3.60E-13	3.90E-18	0.0009	1.30E-10
End	451	557	574	469	558	454	575	557	84
Start	384	509	511	375	518	388	512	514	-
Chain ID	<	<	<	¥	A				Y
PDB ID	lekz	lekz	lekz	1qu6	lqu6	Istu	1sta	1stu	167f
SEQ ID NO:	664	564	664	664	999	664	664	999	599

PDB TD	ਹ <u>ੋ</u>	Chain ID	Start	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
									D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	PROTEIN-RNA COMPLEX, GENE REGULATION/RNA
lcvj H			10	88	7.80E-11	1.03	-		POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*3); CHAIN: M, N, O, P, Q, R, S, T;	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA
1d9a	<		6	88	1.30E-11	0.84	66:0		HU ANTIGEN C; CHAIN: A;	RNA BINDING PROTEIN RNA- BINDING DOMAIN
, L£1	<		8	68	2.60E-11	0.56	0.64		NUCLEOLIN RBD1; CHAIN: A:	STRUCTURAL PROTEIN PROTEIN C23; RNP, RBD, RRM, RNA BINDING DOMAIN, NUCLEOLUS
lsx!			9	88	I.30E-11	0.72	8.0		RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND RNA-BINDING DOMAIN 1SXL 3 (RBD-2), RESIDUES 199 - 294 PLUS N-TERMINAL MET) 1SXL 4 (NMR, 17 STRUCTURES) 1SXL 5	
2sxl			6	88	2.60E-11	1.18	_		SEX-LETHAL PROTEIN; CHAIN: NULL;	RNA-BINDING DOMAIN RNA- BINDING DOMAIN, ALTERNATIVE SPLICING
2u2f	₹		6	88	7.80E-11	0.89	_		SPLICING FACTOR U2AF 65 KD SUBUNIT; CHAIN: A;	RNA-BINDING PROTEIN SPLICING, U2 SNRNP, RBD, RNA-BINDING PROTEIN
2up1	V		2	88	6.50E-11	0.87	-	·	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1
lcrz	<		126	269	0.0013	0.18	9.0		TOLB PROTEIN; CHAIN: A;	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD
lcrz.	<		77	217	3.90E-06	0.21	0.39		TOLB PROTEIN; CHAIN: A;	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND

PDB CI	5 ~	Chain ID	Start	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
										TIOLITED WITH THE
	-									ALPHA/BETA FOLD
lerj		A	-	315	1.80Ē-59	0.2	0.84		TRANSCRIPTIONAL REPRESSOR TUP!; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
lerj	 	A	30	373	3.60E-69	0.24	69:0		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
lgot	 	В	_	172	7.20E-73	0.59	-		GT-ÁLPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
lgot		В	8	420	1.10E-57	0.27	-0.02		GT-ALPHA/GI-ALPHA CHIMERA, CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA: CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
lgot		В	29	372	1.10E-57			91.76	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
lgot		В	71	370	5.40E-56	0.56	8.0		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION
	\dashv									
140s	_	V	9	194	7.80E-14	0.52	-0.18		NICOTINATE	TRANSFERASE DINUCLEOTIDE-

		7										<u> </u>										7	-						T	
PDB annotation	BINDING MOTIF, PHOSPHORIBOSYL TRANSFERASE		TRANSCRIPTION HELIX-BUNDLE	TRANSCRIPTION HELIX-BUNDLE																				TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR	BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGIT ATION				LIGASE CBL, UBCH7, ZAP-70, E2,	UBIQUITIN, E3, PHOSPHORYLATION,
Coumpound	MONONUCLEOTIDE:5,6- CHAIN: A;		TRANSCRIPTION ELONGATION FACTOR S-II; CHAIN: A;	TRANSCRIPTION ELONGATION FACTOR S-II;	CHAIN: A;	TRANSCRIPTION	TEGULATION TO THE STATE OF THE	I KANSCKIP I JONAL	(TEIIS, NUCLEIC-ACID 1TFI 3	BINDING DOMAIN) (NMR, 12	TP ANSCRIPTION	REGULATION	TRANSCRIPTIONAL	ELONGATION FACTOR SII	PINDING DOMAIN ONAB 13	STRUCTURES) 1TFI 4	TRANSCRIPTION	REGULATION	TRANSCRIPTIONAL	ELONGATION FACTOR SII	(TFIIS, NUCLEIC-ACID LIFE)	STRUCTURES) 1TF1 4		TRANSCRIPTION FACTOR PML; CHAIN: NULL;		VIRUS EQUINE HERPES	VIRUS-1 (C3HC4, OR RING	DOMAIN) ICHC 3 (NMR, 1	SIGNAL TRANSDUCTION	PROTEIN CBL; CHAIN: A;
SeqFold score											75.34	+ 7.67												-						
PMF			0.24	0.21		1											-							0.01		9.0			0.45	
Verify score			-0.43	-0.03		0.24											0.26					_		-0.65		-0.48			-0.62	
PSI- BLAST			1.00E-18	1.10E-06		1.80E-21					1 205-21	17-705-1					1.30E-21							1.40E-06		1.80E-13			9.10E-11	
End			64	09		257					257	, Ç					257							342		352			351	
Start				4		211					212	717					212							300		302			277	
Chain ID			<	Y																									4	
PDB ID			1 c 00	1000		146					الم]					111							Ibor		lchc			1fbv	
SEQ S			671	671		671					129				_		671							672		672			672	

PDB annotation	2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	APOPTOSIS INHIBITOR OF APOPTOSIS (IAP), NMR STRUCTURE, BACULOVIRAL 2 IAP REPEAT (BIR), ZINC BINDING DOMAIN	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	DNA-BINDING PROTEIN V(D)) RECOMBINATION ACTIVATING PROTEIN I; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN	PEPTIDE SYNTHETASE GRSA; PEPTIDE SYNTHETASE, GRSA, ADENYLATE FORMING	PEPTIDE SYNTHETASE GRSA; PEPTIDE SYNTHETASE, GRSA, ADENYLATE FORMING	OXIDOREDUCTASE OXIDOREDUCTASE, MONOOXYGENASE, PHOTOPROTEIN, LUMINESCENCE
Coumpound	ZAP-70 PEPTIDE: CHAIN: B: UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7, CHAIN: C;	SIGNAL TRANSDUCTION PROTEIN CBL, CHAIN: A; ZAP-70 PEPTIDE, CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7, CHAIN: C;	INHIBITOR OF APOPTOSIS PROTEIN (2MIHB/C-IAP-1); CHAIN: A;	RAGI; CHAIN: NULL;	RAGI; CHAIN: NULL;	GRAMICIDIN SYNTHETASE 1; CHAIN: A, B; PHENYLALANINE; CHAIN: C. D:	GRAMICIDIN SYNTHETASE 1; CHAIN: A, B; PHENYLALANINE; CHAIN: C, D;	LUCIFERASE; CHAIN: NULL;
SeqFold						158.05		173.59
PMF		0.55	0.37	0.8	0.84		-	
Verify score		-0.4	-0.73	0.34	11.0-		9.0	
PSI- BLAST		3.60E-13	0.0027	2.60E-09	7.20E-09	0	0	3.60E-93
End		349	270	344	342	576	571	574
Start		301	227	267	302	24	39	22
Chain ID		∢	V			V V	4	
PDB ID		1fbv	1qbh	1rmd	Irmd	larn u	lam u	I Ci
SEQ FD		672	672	672	672	673	673	673

PDB annotation	ULL; OXIDOREDUCTASE OXIDOREDUCTASE, MONOOXYGENASE, PHOTOPROTEIN, LUMINESCENCE		TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO-ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION		V; CELL MOTILITY PROTEIN MSP; CYTOSKELETAL PROTEIN, SPERM, CELL MOTILITY PROTEIN	V: CELL MOTILITY PROTEIN MSP: CYTOSKELETAL PROTEIN, SPERM, CELL MOTILITY PROTEIN	A, B, ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	E SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	SMALL GTPASE KARYOPHERIN
Coumpound	LUCIFERASE: CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	MAJOR SPERM PROTEIN; CHAIN: A, B;	MAJOR SPERM PROTEIN; CHAIN: A. B;	SYNTAXIN-1A; CHAIN: A, B, C;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	RAN; CHAIN: A, C;
SeqFold score										
PMF score	-	0.01	0.01	0.01	0.01	0.07	0.07	-0.19	-0.02	90.0
Verify score	0.77	0.21	0.21	0.21	0.21	-0.05	-0.05	0.12	0.16	0.02
PSI- BLAST	3.60E-93	0.0031	0.00054	0.0031	0.00054	7.80E-06	7.80E-06	7.80E-09	1.10E-51	5.40E-07
End	571	192	192	192	192	125	125	286	377	119
Start	36	144	144	144	144	34	34	165		22
Chain ID						V	<	4	¥	æ
PDB ID	llci	1bor	lbor	1bor	1 bor	Imsp	lmsp	[cz3]	1b3u	1 ibr
SEQ ID NO:	673	675	675	919	929	629	089	189	682	682

SEQ ID	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
2									CHAIN: B, D;	TRANSPORT RECEPTOR
682	1qbk	æ	2	400	3.60E-37	0.11	0.07		KARYOPHERIN BETA2; CHAIN: B; RAN; CHAIN: C;	NUCLEAR TRANSPORT PROTEIN COMPLEX HEAT REPEATS, NUCLEAR TRANSPORT PROTEIN COMPLEX
682	3bct		32	363	1.30E-07	-0.18	60.0		BETA-CATENIN; CHAIN: NULL;	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON
683	1e3h	⋖	-	273	7.20E-58	0.55	_		GUANOSINE PENTAPHOSPHATE SYNTHETASE; CHAIN: A;	POLYRIBONUCLEOTIDE TRANSFERASE POLYNUCLEOTIDE PHOSPHORYLASE, GUANOSINE POLYRIBONUCLEOTIDE TRANSFERASE, ATP-GTP 2 DIPHOSPHOTRANSFERASE, RNA PROCESSING, RNA DEGRADATION
683	Гезр	∢		273	7.20E-58	0.62	_		GUANOSINE PENTAPHOSPHATE SYNTHETASE; CHAIN: A;	POLYRIBONUCLEOTIDE TRANSFERASE POLYNUCLEOTIDE PHOSPHORYLASE, GUANOSINE POLYRIBONUCLEOTIDE TRANSFERASE, ATP-GTP DIPHOSPHOTRANSFERASE, 2 RNA PROCESSING, RNA DEGRADATION
684	le5d	⋖	89	161	3.60E-13	0.03	90.0		RUBREDOXIN:OXYGEN OXIDOREDUCTASE; CHAIN: A, B	OXIDOREDUCTASE OXIDOREDUCTASE, OXYGENREDUCTASE, DIIRON- CENTRE, 2 FLAVOPROTEINS, LACTAMASE-FOLD
684	Isml	Y	34	198	1.30E-18	0.29	0.04		PENICILLINASE; CHAIN: A;	HYDROLASE METALLO-BETA- LACTAMASE, ANTIBIOTIC RESISTANCE, BINUCLEAR 2 ZINC, HYDROLASE
684	1sml	¥	498	558	0.00052	-0.22	0.09		PENICILLINASE; CHAIN: A;	HYDROLASE METALLO-BETA- LACTAMASE, ANTIBIOTIC RESISTANCE, BINUCLEAR 2 ZINC, HYDROLASE
684	2bc2	<	45	187	3.60E-10	0.05	-0.05		METALLO BETA- LACTAMASE II; CHAIN: A, B;	HYDROLASE HYDROLASE, BETA- LACTAMASE, ANTIBIOTIC, METALLOENZYME
989	lquq	٧	28	181	3.60E-38	0.01	0.23		REPLICATION PROTEIN A 32	DNA-BINDING PROTEIN RPA, OB-

	Chain ED	Start	End AA	PSI- BLAST	Verify score	PMF	SeqFold	Соитропи	PDB annotation
- }								KD SUBUNIT; CHAIN: A. C. REPLICATION PROTEIN A 14 KD SUBUNIT; CHAIN: B, D;	FOLD, SSDNA-BINDING, DNA- BINDING PROTEIN
∢		62	147	3.60E-13	-0.28	0.11		CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A:	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC THIOREDOXIN
1		09	142	3.60E-07	0.08	0.09		THIOREDOXIN: CHAIN: NULL;	OXIDOREDUCTASE DIMER, THIOREDOXIN, X-RAY CRYSTALLOGRAPHY, OXIDOREDUCTASE
_	A	09	143	3.60E-11	-0.31	0.01		THIOREDOXIN M; CHAIN: A, B;	ELECTRON TRANSPORT ELECTRON TRANSPORT
`	V	70	127	5.40E-07	-0.87	0.18		TRYPAREDOXIN-I; CHAIN: A;	TRYPAREDOXIN TRYX-I; TRYPAREDOXIN, CRITHIDIA FASCICULATA, THIOREDOXIN, 2 TRYPANOSOME, ANOMALOUS DISPERSION, OXIDATIVE STRESS, 3 OXIDOREDUCTASE
-	4	75	181	1.30E-12	0.32	0.34		TRYPAREDOXIN-I; CHAIN: A;	TRYPAREDOXIN TRYX-I; TRYPAREDOXIN, CRITHIDIA FASCICULATA, THIOREDOXIN, 2 TRYPANOSOME, ANOMALOUS DISPERSION, OXIDATIVE STRESS, 3 OXIDOREDITCTASE
	A	48	233	5.40E-40	0.06	-0.14		HUMAN THIOREDOXIN PEROXIDASE-B; CHAIN: A, B, C, D, E, F, G, H, I, J;	PEROXIDASE 2-CYS PEROXIREDOXIN, CALPROMOTIN PEROXIDASE, PEROXIREDOXIN, SULPHINIC ACID, THIOREDOXIN
	¥	48	210	1.60E-40	0.1	0.09		THIOREDOXIN PEROXIDASE 2; CHAIN: A, B;	OXIDOREDUCTASE HEME-BINDING PROTEIN 23 KD, HBP23; THIOREDOXIN FOLD, OXIDOREDUCTASE
7 1	V	78	187	1.30E-06	-0.66	0.11		THIOREDOXIN; CHAIN: A;	ELECTRON TRANSPORT ALPHA/BETA OPEN-TWISTED PROTEIN, THIOL- DISULFIDE
	æ	62	150	1.60E-12	-0.36	0.03		DNA POLYMERASE; CHAIN: A; THIOREDOXIN; CHAIN: B; DNA; CHAIN: P, T;	T7 DNA POLYMERASE, DNA REPLICATION, NUCLEOTIDYL 2 TRANSFERASE, SEQUENCING, THIOREDOXIN, PROCESSIVITY FACTOR, 3 COMPLEX

PDB Ch	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
		¥	¥ *	DLASI	score	score.	31008		
	 								(HYDROLASE/ELECTRON TRANSPORT/DNA)
<		62	150	1.60E-12	-0.1	0.04		ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2 2TRXA 3	
		318	363	7.20E-13	-0.33	0.48		VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC3 (NMR, 1 STRUCTURE) ICHC 4	
<		298	363	1.20E-10	-0.58	0.21		SIGNAL TRANSDUCTION PROTEIN CBL, CHAIN: A; ZAP-70 PEPTIDE, CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	LIGASE CBL, UBCH1, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
∢		319	368	1.80E-06	-0.74	0.21		SIGNAL TRANSDUCTION PROTEIN CBL, CHAIN: A; ZAP-70 PEPTIDE, CHAIN: B; UBIQUITIN-CONJUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	LIGASE CBL, UBCH7, ZAP-70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,
Irmd		297	363	1.00E-13	-0.07	0.94		RAGI; CHAIN: NULL;	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN
lrınd		319	368	5.40E-06	-0.24	0.49		RAG1; CHAIN: NULL;	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA- BINDING PROTEIN
4		4	307	5.40E-42	0.66	0.11		TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTION INHIBITOR BETA- PROPELLER
4		71	369	9.00E-51	0.52	0.29		TRANSCRIPTIONAL	TRANSCRIPTION INHIBITOR BETA-

PDB annotation	A, PROPELLER	COMPLEX (GTP-BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT: GAMMA!, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA- A, PROPELLER	TRANSCRIPTION INHIBITOR BETA- A, PROPELLER	TRANSCRIPTION INHIBITOR BETA- A, PROPELLER	TRANSCRIPTION INHIBITOR BETA-
Coumpound	REPRESSOR TUP1; CHAIN: A, B. C.	GT-ÁLPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A, GT- BETA; CHAIN: B, GT- GAMMA; CHAIN: G;	TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A. B. C.	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B. C.	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL
SeqFold score		70.14							
PMF score			0.16	-0.17	0.29	0.94	_	0.99	0.16
Verify score			0.39	0.26	0.2	0.12	0.46	0.01	0.27
PSI- BLAST		3.60E-57	3.60E-57	1.40E-36	9.00E-08	5.40E-78	1.80E-65	9.00E-57	1.40E-52
End		376	370	304	377	416	313	442	224
Start		28	39	s	133	801	81	195	5
Chain ID		В	<u>ш</u>	മ	¥.	¥	4	<	4
PDB CD		lgot	1got	lgot	1crz	lerj	1erj	lerj	lerj
SEQ	2	693	693	693	694	694	694	694	694

PDB annotation		COMPLEX (GTP-BINDING/TRANSDUCER) BETA!, BINDING/TRANSDUCER) BETA!, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN.	HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA	SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBINIT:	GAMMAI, TRANSDUCIN GAMMA	SUBDINIT, COMPLEX (OTF-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL	TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1,	TRANSDUCIN BETA SUBUNIT;	SUBUNIT; COMPLEX (GTP-	BINDING/TKAN3DOCEN), OTROTEILY, HETEROTRIMER 2 SIGNAL TRANSDICTION	COMPLEX (GTP-	TRANSDUCIN BETA SUBUNIT;	GAMMAI, TRANSDUCIN GAMMA	SUBUNIT; COMPLEX (GIP-BINDING/TRANSDUCER), G PROTEIN,	HETEROTRIMER 2 SIGNAL TRANSDUCTION
Coumpound	B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- RETA: CHAIN: B: GT-	GAMMA; CHAIN: G;			GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT-	BETA, CHAIN: B; GT-	() () () () () () () () () () () () () (GT-ALPHA/GI-ALPHA	BETA; CHAIN: B; GT-	GAMMA; CHAIN: G;		
SeqFold score				105.8														
PMF		_				0.21				0.22				_				
Verify score		0.54				0.26				0.36				69.0				
PSI- BLAST		5.40E-64		7.20E-81		3.60E-50				1.80E-53				7.20E-81				
End AA		393		353		443				267				353	·			
Start		103		12		061				4				26				
Chain ID		В		В		В				В				В				
PDB TD		lgot		1got		1got				lgot				lgot		_		-
SEQ ID NO:		694		694		694				694				694				

SEQ	PDB	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
≘ ÿ	e	e	ΥΥ	VV	BLAST	score	score	score		
										Very Control of Control
269	lalh	4	200	282	1.30E-31			84.99	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OFFIGONTICLEOTIDE	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
					_				BINDING SITE; CHAIN: B, C,	
269	lalh	<	228	309	3.60E-27	0.18	_		QGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
									PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC FINGER DNA-RINDING PROTEIN
									BINDING SITE; CHAIN: B, C;	
269	laih	<	228	337	1.30E-31	-0.13	0.03		QGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
									PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC
									OLIGONUCLEOTIDE	FINGER, DNA-BINDING PROLEIN
203	1.51	\ \	341	121	7 205 30	0 03	0.81		OGGR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
/60	i a i	<	1 + 6	174	1.20E-30	6.0	0.01		PEPTIDE: CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC
									OLIGONÚCLEOTIDE	FINGER, DNA-BINDING PROTEIN
									BINDING SITE; CHAIN: B, C;	
269	lalh	٧	397	480	1.80E-28	0	69.0		QGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
									PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC
		_							OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C.	FINGER, DNA-BINDING PROTEIN
269	1811	4	40	86	\$ 40E-10	0.13	-0.17		OGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
		•	<u>:</u>	· _	}	!	;		PEPTIDE: CHAIN: A, DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC
									OLIGONUCLEOTIDE	FINGER, DNA-BINDING PROTEIN
									BINDING SITE; CHAIN: B, C;	
269	lalh	Ą	484	554	5.40E-28	0.14	69.0		QGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
			_						PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC
									OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	FINGER, DNA-BINDING PROTEIN
169	laih	4	74	154	3.60E-26	-0.1	0.39		QGSR ZINC FINGER	COMPLEX (ZINC FINGER/DNA)
									PEPTIDE; CHAIN: A; DUPLEX	COMPLEX (ZINC FINGER/DNA), ZINC
				_					OLIGONUCLEOTIDE	FINGER, DNA-BINDING PROTEIN
									BINDING SITE; CHAIN: B, C;	
169	1550		514	264	6.50E-11	-0.35	60:0		DNA-BINDING PROTEIN	
									HUMAN ENHANCER-	
									MITANT WITH CVC 11 18BO	
						_			3 REPLACED BY ABU	
									(C11ABU) (NMR, 60	
									STRUCTURES) 1BBO 4	
697	Ime	၁	101	168	1.80E-36	-0.09	0.45		DNA; CHAIN: A, B, D, E;	COMPLEX (ZINC FINGER/DNA) ZINC

SEQ ID NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
	٨								CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
269	Ime y	U .	102	224	1.30E-16	-0.26	0.16		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
269	Ime y	ပ	156	252	9.00E-35	-0.15	8.0		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
697	, y	O	199	280	5.40E-47	0.31			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
697	1me y	O	661	281	5.40E-47			97.32	DNA: CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
697	Ime y	ပ	227	309	3.60E-46	0.27	96.0		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
697	lme y	U	227	337	1.30E-33	-0.23	0.22		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
697	Lme Y	O	283	365	3.60E-47	0.21	0.95		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	COMPLEX (ZINC FINGERIDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
697	y me	ပ	312	393	1.40E-49	0.41	0.99		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA

d PDB annotation			CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)			CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	L	FINGER FINGER, PROTEIN-UNA	 (ZINC FINGER/DNA)	D, E; COMPLEX (ZINC FINGER/DNA) ZINC						(ZINC FINGER/DNA)				CRISIAL SIRUCIORE, COMPLEX (ZINC FINGER/DNA)		_		CRYSTAL STRUCTURE, COMPLEX			_
Coumpound	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN: CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER		DNA; CHAIN: A, B, D, E;	PROTEIN: CHAIN: C. F. G.		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	-	DNA. CHAIN. A D	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G,
SeqFold score					-															. <u>.</u>							
PMF score		0.95	<u> </u>	0.88			0.04			0.62			0				98.0				-				000	0.70	
Verify score		0.09		0			-0.68			0.03			-0.13	_			0.22				0.31				01.0	0.70	
PSI- BLAST		1.30E-48		1.80E-48			1.30E-21			3.60E-47			1.60E-35				3.60E-48			-	3.60E-50				1 100 44	1.105-44	
End		422		451			208			480			126				208				536			_	667) CC	
Start AA		340		368		_	368			396			42				425				455				402	6	
Chain ID		၁		2			J			၁			2				C				O	_			(رر	
PDB ID		L mc		Imc	۲.		E E	^		Imc	`		Ime	λ.			Ime	γ			Ime	>) 	`
SEQ D	1	269		269			269			269			269				269				169				207	/60	

PDB annotation	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC	NTERACTION, PROTEIN DESIGN, 2	CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TFIIIA; 5S GENE;	TRANSCRIPTION EACTOR SERVING	GENE, DNA BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	REGULATION/DNA). RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	KEGULATION/DNA) COMPLEX	PEGIII ATIONONA) BNA	POLYMERASE III. 2 TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX	REGILLATION/DNA) RNA	POLYMERASE III, 2 TRANSCRIPTION
Coumpound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN, CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E;	PROTEIN: CHAIN: C, F, G;		TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 5S RNA	GENE; CHAIN: E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	(1, 2, 5, 5, 1)			TFIIIA; CHAIN: A, D; 5S	KIBUSUMAL KNA GENE;	(1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1			TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C. E. F;	
SeqFold score															107.54													
PMF		0.58		0.94		-0.13			9.0											0.92					0.75			
Verify		-0.2		0.53		0.3			-0.1											-0.08					-0.02			
PSI- BLAST		1.80E-43		5.40E-13		1.10E-09			1.10E-19						2.60E-50					3.60E-37					3.60E-34			
End		154		536		86			417						366]	346					461			
Start AA		73		509		7.1			341						199				3	200					313			
Chain ID		၁		9		Ð			∢						∢					<					<			
PDB ID		Ime y		lme y		, me			113					87.	1116			-	2	911					931			
SEQ TO NO:		269		. 269		169			269					,	/69				į	/69					269			

PMF SeqFold score score 0.23
0.72
0.01
69.0
0.51
0.95
0.28

PDB annotation	REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)
Coumpound	ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SeqFold score				-		83.84
PMF score		0.7	66.0	_	0.51	
Verify		-0.3	-0.03	1.0-	0	
PSI- BLAST		1.80E-33	9.00E-35	2.60E-28	3.60E-32	1.80E-34
End		365	422	450	508	537
Start		263	320	345	404	427
Chain ID		ن ن	ပ	U	ပ	ပ
PDB ID		pqnI	lubd	1 ubd	pqnI	1 ubd
SEQ ID NO:		769	169	269	269	697

PDB annotation			. Ý	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADR1, ZINC FINGER, NMR	HON 4	GLI1; COMPLEX (DNA-BINDING C, C, PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
Coumpound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY I; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ADR1; CHAIN: NULL;	COMPLEX(TRANSCRIPTION REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;
SeqFold score							
PMF score	0.88	0.46	0.35	0.07	0.31	-0.12	0.45
Verify	90.0	-0.05	-0.25	-0.45	-0.15	0.26	-0.08
PSI- BLAST	1.80E-34	3.60E-32	1.30E-26	1.30E-18	2.60E-14	1.30E-15	7.20E-26
End	536	558	154	253	260	564	279
Start	433	463	69	78	512	808	101
Chain ID	O	U	S	၁		A	₹
PDB ID	lubd	Pdul	lubd	1ubd	2adr	24гр	2gli
SEQ ID NO:	697	697	697	697	269	697	269

PDB Chain Start ID ID AA	Start AA			End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
2gii A 102 282 6.5	102 282	282		6.5	6.50E-40	-0.17	66.0		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 198 308 7.20	198 308	308		7.20	7.20E-28	0.14	_		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gii A 200 394 5.2C	200 394	394	 -	5.20	5.20E-48	-0.24	0.25		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 235 367 1.8	235 367	367		1.80	1.80E-32	90.06	66.0		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
28li A 255 397 5.20	255 397	397	-	5.20	5.20E-48			89.2	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 292 422 5.40E-35	292 422	422	<u> </u>	5.401	3-35	-0.02	0.78		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 320 449 3.60E-34	320 449	449		3.601	9-34	0.04	0.86		ZINC FINGER PROTEIN GLII; CHAIN: A: DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER. COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 404 535 5.40	404 535	535		5.40	5.40E-33	0.2	0.92		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 433 558 1.60	.433 558	558		1.60	1.60E-31	0.02	0.22		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 66 153 1.1	66 153	153		=	1.10E-24	-0.05	0.03		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
2gli A 77 253 3.9	77 253	253	Н	3.9	3.90E-30	-0.45	0.13		ZINC FINGER PROTEIN GL11;	COMPLEX (DNA-BINDING

SEQ TO	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									CHAIN: A; DNA; CHAIN: C, D;	PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
869	1edh	∢	101	302	7.20E-32	1.0-	0.63		E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
869	ledh	V	74	300	7.20E-32			62.4	E-CADHERIN; CHAIN: A, B;	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12, CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN
869	1ncj	¥	9	178	1.40E-34	0.04	-0.12		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
869	lncj	Ą	70	302	1.80E-32			64.18	N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
869	lncj	¥	74	302	1.80E-32	-0.03	0.16		N-CADHERIN; CHAIN: A;	CELL ADHESION PROTEIN CELL ADHESION PROTEIN
869	Isuh		23	62	1.30E-06	-0.22	0.04		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
869	1suh		74	182	1.30E-08	-0.08	0.01		EPITHELIAL CADHERIN; CHAIN: NULL;	CELL ADHESION UVOMORULIN; CADHERIN, CALCIUM BINDING, CELL ADHESION
701	1a7j		128	281	1.30E-13	0.17	9.0		PHOSPHORIBULOKINASE; CHAIN: NULL:	TRANSFERASE TRANSFERASE, KINASE, CALVIN CYCLE
701	187j		95	366	1.30E-13			6.99	PHOSPHORIBULOKINASE; CHAIN: NULL;	TRANSFERASE TRANSFERASE, KINASE, CALVIN CYCLE
701	1bd3	Q	306	532	2.60E-73			191.89	URACIL PHOSPHORIBOSYLTRANSFE RASE: CHAIN: D, C, B, A;	TRANSFERASE UPRTASE; TRANSFERASE, GLYCOSYLTRANSFERASE, UPRTASE
701	1bd3	Ω	322	532	3.60E-44	0.84	-		URACIL PHOSPHORIBOSYLTRANSFE RASE: CHAIN: D. C. B. A:	TRANSFERASE UPRTASE; TRANSFERASE, GLYCOSYLTRANSFERASE, UPRTASE
701	1643	Q	324	532	2.60E-73	0.76	-		URACIL PHOSPHORIBOSYLTRANSFE RASE: CHAIN: D. C. B. A:	TRANSFERASE UPRTASE; TRANSFERASE, GLYCOSYLTRANSFERASE, UPRTASE
701	lesm	Ą	75	294	1.80E-35	0.43	96.0		PANTOTHENATE KINASE,	TRANSFERASE PANK; PROTEIN-

SEQ ID	PDB	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
Ö									CHAIN: A, B, C, D;	INHIBITOR COMPLEX
101	lesm	4	16	303	6.50E-67	0.55	66.0		PANTOTHENATE KINASE; CHAIN: A, B, C, D;	TRANSFERASE PANK; PROTEIN- INHIBITOR COMPLEX
101	lukz		94	302	1.20E-15	0.19	0.54		TRANSFERASE URIDYLATE KINASE (E.C.2.7.4) COMPLEXED WITH ADP AND AMP I UKZ 3	
101	1zin		105	302	0.0078	-0.33	0.21		ADENYLATE KINASE; CHAIN: NULL;	PHOSPHOTRANSFERASE ADK; PHOSPHOTRANSFERASE, ZINC FINGER
101	3tmk	C	96	303	1.30E-23	-0.38	0.03		THYMIDYLATE KINASE; CHAIN: A, B, C, D, E, F, G, H;	KINASE KINASE, PHOSPHOTRANSFERASE
702	1bq0		∞	84	1.30E-31			54.61	DNAJ; CHAIN: NULL;	CHAPERONE, HSP40; CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK
702	1bq0		6	84	1.30E-31	0.52	0.52		DNAJ; CHAIN: NULL;	CHAPERONE HSP40, CHAPERONE, HEAT SHOCK, PROTEIN FOLDING, DNAK
702	lcvj	<	182	255	3.60E-14	0.07	69:0		POLYDENYLATE BINDING PROTEIN I; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA
702	lcvj	tı.	781	259	1.80E-12	0.13	0.46		POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, Q, R, S, T;	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA
702	lcvj	н	182	259	1.80E-12	0.03	0.46		POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP*AP *AP*AP*AP*A)-3'); CHAIN: M, N, O, P, O, R, S, T;	GENE REGULATION/RNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN-RNA COMPLEX, GENE REGULATION/RNA
702	148z	4	185	251	1.80E-12	0.15	0.93		HU ANTIGEN C; CHAIN: A;	RNA BINDING PROTEIN RNA- BINDING DOMAIN
702	lez3	<	77	186	1.30E-09	0.14	-0.13		SYNTAXIN-1A; CHAIN: A, B, C;	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35

PDB ID		Chrin ID	Start AA	End AA	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
										KDA PROTEIN, P35A, THREE HELIX BUNDLE
lha l	}		182	249	1.30E-22	0.33	0.19		HNRNP A1; CHAIN: NULL;	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN AI, NUCLEAR PROTEIN, HNRNP, RBD, RRM, RNP, RNA BINDING, 2 RIBONUCLEOPROTEIN
1141		<	182	249	5.40E-16	0.12	0.51		HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN DO; CHAIN: A;	RNA BINDING PROTEIN RNA- BINDING DOMAIN
l hdj			=	78	2.60E-19	0.57	_		HUMAN HSP40; CHAIN: NULL;	MOLECULAR CHAPERONE HDI-1; MOLECULAR CHAPERONE
1hdj			8	84	1.80E-28			56.45	HUMAN HSP40; CHAIN: NULL;	MOLECULAR CHAPERONE HDI-1; MOLECULAR CHAPERONE
!hdj			8	84	1.80E-28	10.0	0.47		HUMAN HSP40; CHAIN: NULL;	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE
2mss		∀	182	249	7.20E-17	0.42	0.18		MUSASHII; CHAIN: A;	RNA BINDING PROTEIN RNA- BINDING DOMAIN
2up1		Y	165	255	5.40E-26	0.16	0		HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	COMPLEX (RIBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1
2up1		· V	182	259	1.30E-18	-0.11	0.05		HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12-NUCLEOTIDE SINGLE-STRANDED TELOMETRIC DNA; CHAIN: B;	COMPLEX (RJBONUCLEOPROTEIN/DNA) HNRNP A1, UP1; COMPLEX (RJBONUCLEOPROTEIN/DNA), HETEROGENEOUS NUCLEAR 2 RJBONUCLEOPROTEIN A1
lez3	+	V V	468	575	6.50E-06	0.15	0.11		SYNTAXIN-1A; CHAIN: A, B, C;	ENDOCYTOSIS/EXOCYTOSIS SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE
lawc	+	В	137	292	1.80E-36			63.45	GA BINDING PROTEIN	COMPLEX (TRANSCRIPTION

SeqFold Coumpound PDB annotation score	ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E; REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E; REGULATION/DNA), DNA-BINDING, 2 REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	GA BINDING PROTEIN ALPHA: CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E; REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION (TRANSCRIPTION) REGULATION/DNA). DNA-BINDING, 2 REGULATION/DNA). DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E; REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX CHAIN: B; DNA; CHAIN: D, E; REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR		ANKYRIN MOTTE ANKYRIN MOTTE P19INK4D CDK4/6 TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, CHAIN: NULL;
PMF Se score			-	0.86	1	
Verify score		0.84	0.49	0.51	0.85	0.72
PSI- BLAST		1.30E-30	1.80E-36	7.20E-33	3.60E-31 3.60E-31	9.00E-31
End		273	323	351	298	320
Start AA		169	175	209	144	178
Chain ID		В	B	В		
PDB ID		lawc	lawc	lawc	15d8 15d8	1bd8
SEQ NO:		706	706	706	706	902

PDB annotation	SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEINACINA SE) INHIBITOR	PROTEIN, CYCLIN-DEPENDENT KINASE CELL CYCLE 2 CONTROL	ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR	PROTEIN, CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2 CONTROL,	ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR	PROTEIN/KINASE) INHIBITOR	PROTEIN, CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2 CONTROL,	ALPHA/BETA, COMPLEX (INHIBITOR	HODAKONE/CROWTH FACTOR P18	INKAC: CELL CYCLE INHIBITOR	P18INK4C, TUMOR, SUPPRESSOR,	CYCLIN- 2 DEPENDENT KINASE,	HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18-	PIRINK 4C TITMOR SUPPRESSOR	CYCLIN- 2 DEPENDENT KINASE	HORMONE/GROWTH FACTOR	SIGNALING PROTEIN HELIX-TURN-	HELIX, ANKYRIN REPEAT	CELL CYCLE NHIBITOR P18- INK4C(INK6); CELL CYCLE
Coumpound	INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;		CYCLIN-DEPENDENT KINASE 6: CHAIN: 4:	P19INK4D; CHAIN: B;		CYCLIN-DEPENDENT	P19INK4D; CHAIN: B;			CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;	P19INK4D, CHAIN: B;			CVCI IN DEPENDENT	KINASE 6 INHIBITOR:	CHAIN: A:			CYCLIN-DEPENDENT	CHAIN: A:	•		CYCLIN-DEPENDENT	KINASE 4 INHIBITOR B; CHAIN: A:	CYCLIN-DEPENDENT KINASE 6 INHIBITOR;
SeqFold score		63.41														74 38	00:1										68.42
PMF				_			96.0				-0.13				,						0.95				_		
Verify				0.71			0.65				0.24										0.61				0.55		
PSI- BLAST		1.10E-30		1.80E-30			1.10E-30				3.60E-26					9 00F-34					9.00E-34				1.30E-28		5.40E-33
End		297		293		·	315				224					302	1				329				273		296
Start AA		144		145			178				57					134	-				175				121		138
Chain ID		В		В			В	_			В					4	:				<				٧		¥
PDB ID		1blx		1blx			1blx				1blx		_			11419				1	1bu9	_			1d9s		l ihb
SEQ ID NO:		706		706			706				902			_		706	2			1	706				706		706

SEQ ID NO:	PDB ID	Chain ID	Start AA	End AA	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									CHAIN: A, B;	INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR
706	lihb	<	175	328	5.40E-33	0.88	0.99		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR
706	Likn	Q	82	290	7.20E-41	0.26	0.96		NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	TRANSCRIPTION FACTOR P65, P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX
706	l ikn	Q	95	307	7.20E-41			74.07	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX
706	lmy o		691	288	2.60E-32			66'09	MYOTROPHIN; CHAIN: NULL	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK-REPEAT
706	1my 0		172	287	2.60E-32	0.32	_		MYOTROPHIN; CHAIN: NULL	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK-REPEAT
706	Infi	шì	12	208	1.10E-36	0	-0.06		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANK YRIN 2 REPEAT HELIX
706	Infi	tr)	136	310	5.40E-36	69:0	-		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; 1-KAPPA-B-ALPHA; CHAIN: E. F:	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX
706	Infi	пĵ	169	341	1.80E-34	0.55	0.46		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F:	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX
706	Infi	tr)	204	414	3.60E-27	0.18	0.18		NF.KAPPA-B P65; CHAIN: A, C; NF.KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX
706	Infi	ம	18	290	3.60E-41	0.26	86.0		NF.KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E. E.	COMPLEX (TRANSCRIPTION REGIANK REPEAT) COMPLEX (TRANSCRIPTION REGULATIONANK REPEAT) ANKYRIN 2 REPEAT HELIX
902	lnfi	ы	95	303	3.60E-41			71.81	NF-KAPPA-B P65; CHAIN: A,	COMPLEX (TRANSCRIPTION

ρ. –	PDB ID	Chain 1D	Start AA	End AA	PSI- BLAST	Verify	PMF score	SeqFold score	Coumpound	PDB annotation
	, 								C; NF-KAPPA-B P50; CHAIN: B, D; 1-KAPPA-B-ALPHA; CHAIN: E, F;	REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX
lycs		B	170	362	1.80E-25			75.09	P53; CHAIN: A; 53BP2; CHAIN: B;	COMPLEX (ANTI- ONCOGENE/ANKYRIN REPEATS) PS3BP2; ANKYRIN REPEATS, SH3, PS3, TUMOR SUPPRESSOR, MULTIGENE 2 FAMILY, NUCLEAR PROTEIN, PHOSPHORYLATION, DISEASE MUTATION, 3 POLYMORPHISM, COMPLEX (ANTI- ONCOGENE/ANKYRIN REPEATS)
1n8y	 		19	336	5.40E-42			87.98	CALSEQUESTRIN; CHAIN: NULL	CALCIUM-BINDING PROTEIN CALSEQUESTRIN, CALCIUM-BINDING PROTEIN, SARCOPLASMIC 2 RETICULUM, RABBIT SKELETAL MUSCLE
1a8y			20	332	5.40E-42	0.15	96.0		CALSEQUESTRIN; CHAIN: NULL	CALCIUM-BINDING PROTEIN CALSEQUESTRIN, CALCIUM-BINDING PROTEIN, SARCOPLASMIC 2 RETICULUM, RABBIT SKELETAL MUSCLE
16jx	-		133	239	1.60E-13	-0.03	0.07		PROTEIN DISULFIDE ISOMERASE; CHAIN: NULL;	ELECTRON TRANSPORT ELECTRON TRANSPORT, REDOX-ACTIVE CENTER, ISOMERASE, 2 ENDOPLASMIC RETICULUM
1dby		<	23	131	5.40E-24			52.47	CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A;	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC THIOREDOXIN
1dby		Y	30	131	5.40E-24	0.3	0.95		CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A:	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC THIOREDOXIN
lerv			14	128	1.10E-23	0.54	86.0		THIOREDOXIN; CHAIN: NULL;	OXIDOREDUCTASE DIMER, THIOREDOXIN, X-RAY CRYSTALLOGRAPHY, OXIDOREDUCTASE
1126		V	27	130	9.00E-25	0.81	_		THIOREDOXIN M; CHAIN: A, B;	ELECTRON TRANSPORT ELECTRON TRANSPORT
는 교			20	134	1.10E-31			82.03	PROTEIN DISULFIDE ISOMERASE; CHAIN: NULL;	ELECTRON TRANSPORT ELECTRON TRANSPORT, REDOX-ACTIVE

	, —					,					
PDB annotation	CENTER, ISOMERASE, 2 ENDOPLASMIC RETICULUM	ELECTRON TRANSPORT ELECTRON TRANSPORT, REDOX-ACTIVE CENTER, ISOMERASE, 2 ENDOPLASMIC RETICULUM	ELECTRON TRANSPORT ALPHA/BETA OPEN-TWISTED PROTEIN, THIOL- DISULFIDE	T7 DNA POLYMERASE, DNA REPLICATION, NUCLEOTIDYL 2 TRANSFERASE, SEQUENCING, THIOREDOXIN, PROCESSIVITY FACTOR, 3 COMPLEX (HYDROLASE/ELECTRON TRANSPORT/DNA)	T7 DNA POLYMERASE, DNA REPLICATION, NUCLEOTIDYL 2 TRANSFERASE, SEQUENCING, THIOREDOXIN, PROCESSIVITY FACTOR, 3 COMPLEX (HYDROLASE/ELECTRON TRANSPORT/DNA)	ELECTRON TRANSPORT THIOREDOXIN 2; ITHX 7 OXIDO- REDUCTASE ITHX 16	ELECTRON TRANSPORT THIOREDOXIN 2; ITHX 1 OXIDO- REDUCTASE ITHX 16	ELECTRON TRANSPORT HTRX, HCHI, CHI; OXIDOREDUCTASE, ELECTRON TRANSPORT			DNA INTEGRATION
Coumpound		PROTEIN DISULFIDE ISOMERASE; CHAIN: NULL;	THIOREDOXIN; CHAIN: A;	DNA POLYMERASE; CHAIN: A; THIOREDOXIN; CHAIN: B; DNA; CHAIN: P, T;	DNA POLYMERASE; CHAIN: A; THIOREDOXIN; CHAIN: B; DNA; CHAIN: P, T;	THIOREDOXIN: ITHX 5 CHAIN: NULL; ITHX 6	THIOREDOXIN; ITHX 5 CHAIN: NULL; ITHX 6	THOREDOXIN H; CHAIN: NULL;	ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2 2TRXA 3	ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2 2TRXA 3	AVIAN SARCOMA VIRUS INTEGRASE; 1ASU 7 CHAIN:
SeqFold score				60.72		55.88			62.2		
PMF		86:0	1		_		-	99'0		_	0.39
Verify score		0.43	0.62		0.39		0.75	0.33		0.39	-0.13
PSI- BLAST		1.10E-31	3.60E-24	1.60E-24	1.60E-24	9.10E-22	9.10E-22	9.00E-23	5.40E-25	5.40E-25	1.80E-25
End		133	130	128	130	131	125	129	131	130	283
Start		21	30	26	27	24	32	25	24	25	130
Chain 1D			¥	В	В				٧	A	
PDB ID		Ime k	lqu *	1t7p	1t7p	1thx	1thx	ltof	2trx	2trx	lasu
SEQ ID NO:		709	709	709	709	709	709	709	709	709	715

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PDB annotation		TRANSFERASE DNA INTEGRATION	TRASFERASE DNA INTEGRATION, TRASFERASE	DNA INTEGRATION DNA	INTEGRATION, AIDS, POLYPROTEIN,	HYDROLASE, 2 ENDONUCLEASE,	POLYNUCLEOTIDYL TRANSFERASE,	DNA BINDING 3 (VIRAL)	TRANSFERASE INTEGRASE, ROUS	SARCOMA VIRUS, HIV, X-RAY	CRYSTALLOGRAPHY, 2 PROTEIN	STRUCTURE, TRANSFERASE	VIRUS/VIRAL PROTEIN INTEGRASE,	ROUS SARCOMA VIRUS, HIV, X-RAY	CRYSTALLOGRAPHY, 2 VIRUS/VIRAL	PROTEIN	TRANSFERASE MIXED BETA-SHEET	SURROUNDED BY ALPHA-HELICES	VIRUS/VIRAL PROTEIN HIV-1	INTEGRASE, POLYNUCLEOTIDYL	TRANSFERASE, DNA-BINDING 2	PROTEIN, DD35E	HYDROLASE DNA INTEGRATION,	INTEGRASE, HIV, HYDROLASE,	ASPARTYL 2 PROTEASE,	ENDONOCEEASE	ANTI-ONCOGENE CELL CYCLE, ANTI-	ONCOGENE, REPEAT, ANK REPEAT	ANTI-ONCOGENE CELL CYCLE, ANTI-	ONCOGENE, REPEAT, ANK REPEAT	COMPLEX (TRANSCRIPTION	REGULATION/DNA) GABPALPHA;	GABPBETA1; COMPLEX	(TRANSCRIPTION	REGULATION/DNA), DNA-BINDING,	NUCLEAR PROTEIN, ETS DOMAIN,	ANKYRIN REPEATS, TRANSCRIPTION	3 FACTOR
Coumpound	NUCL; 1ASU 8	INTEGRASE, CHAIN: A;	INTEGRASE; CHAIN: A;	INTEGRASE; CHAIN: A, B, C;					INTEGRASE; CHAIN: A, B, C,	Ď;			RSV INTEGRASE; CHAIN: A,	B;			AVIAN SARCOMA VIRUS	INTEGRASE; CHAIN: A;	POL POLYPROTEIN, CHAIN:	A, B;			HIV-I INTEGRASE; CHAIN:	A, B, C;			TUMOR SUPPRESSOR	P16INK4A; CHAIN: NULL;	TUMOR SUPPRESSOR	P16INK4A; CHAIN: NULL;	GA BINDING PROTEIN	ALPHA; CHAIN: A; GA	BINDING PROTEIN BETA 1;	CHAIN: B; DNA; CHAIN: D, E;				
SeqFold score																					_												_					
PMF score		0.29	0.21	0.19					0.1				0.13				89.0		0.17				0.39				66.0		_		_							
Verify score		0.21	0.07	0					90:0-				-0.25				80.0		21.0				0.41				0.41		89.0		0.12							
PSI- BLAST		5.40E-26	5.40E-31	5.40E-33					1.30E-29				3.60E-26				3.60E-21		9.00E-26				9.00E-28				1.30E-24		6.50E-24		1.10E-34							00 007
End		297	297	297					337				337				274		297				297				228		161		239							2,5
Start AA		142	142	142					131				139				135		142				142				112		94		112							33
Chain ID		V	<	C					٧				В				∢		∢				<								æ			_				,
PDB ID		1691	169t	1513					100	E			lcla				lcxd		lexd				lqs4				1a5c		laSe		lawc							
SEQ 5 D S		715	715	715					715				715				715		715				715				416		719		719							710

PDB annotation	REGULATIONDNA) GABPALPHA, GABPBETA1, COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (KINAŜE/ANTI- ONCOGENE) CDK6, PI6INK4A, MTS1, CYCLIN DEPENDENT KINAŜE, CYCLIN DEPENDENT KINAŜE, CYCLIN DEPENDENT KINAŜE, CHIBITORY 2 PROTEIN, CDK, INK4, CELL CYCLE, MULTIPLE TUMOR SUPPRESSOR, 3 MTS1, COMPLEX (KINAŜE/ANTI-ONCOGENE) HEADER	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
Coumpound	ALPHA, CHAIN: A: GA BINDING PROTEIN BETA I; CHAIN: B: DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A, MULTIPLE TUMOR SUPPRESSOR; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;			
SeqFold score				68.17				
PMF		-	0.51			-	-	0.36
Verify score		0.32	-0.08		0.77	0.13	0.33	0.21
PSI- BLAST		5.40E-39	1.10E-25	7.80E-35	7.80E-35	1.10E-34	1.40E-25	9.00E-24
End		228	160	233	218	231	228	091
Start AA		78	21	73	84	78	112	21
Chain ID		æ					В	В
PDB ID		Івис	1 pq8	1bd8	8Pq1	8pq1	16i7	1blx
SEQ ID NO:		719	719	719	917	416	719	719

Ē	- m	Chain	Start	End	PSI-	Verify	PMF	SeqFold	Coumpound	PDB annotation
8		<u>e</u>	٧٧	¥	BLAST	score	score	score		!
× Iq1	 	В	44	201	1.30E-35			70.26	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
161x	 	æ	78	218	1.30E-35	0.62	-		CYCLIN-DEPENDENT KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
1blx		В	78	231	3.60E-34	0.39	-		CYCLN-DEPENDENT KINASE 6, CHAIN: A; P19INK4D; CHAIN: B;	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
16u9		∢	18	165	1.10E-25	-0.07	0.12		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C; TUMOR, SUPPRESSOR. CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR
1bu9		K	70	239	1.10E-36			67.41	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18NK4C, TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR
1bu9		¥.	78	233	1.10E-36	0.17	-		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR
149s		¥	101	218	2.60E-29	0.33	-		CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT
149s		∢	112	234	1.10E-25	0.48	66:0		CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT
149s		4	78	197	1.20E-28	0.73	-		CYCLIN-DEPENDENT KINASE 4 INHIBITOR B;	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT

Ith A 18 140E-21 -0.2 0.06 CTCLIN-	0	PDB	Chain	Start	End	-ISA	Verify	PMF	SeqFold	Coumpound	PDB annotation
Itiho A 1 131 140E-21 -0.2 0.06 CYCLIN-DEPENDENT	ЭŠ	<u>e</u>	ല	AA	¥¥	BLAST	score	score	score		
Ith A 1 131 140E-21 -0.2 0.06 CYCLIN-DEPENDENT	П									CHAIN: A;	
Itiho A 18 164 5.40E-25 0.31 0.93 CYCLIN-DEPENDENT	719	lihb	V	1	131	1.40E-21	-0.2	90'0		CYCLIN-DEPENDENT KINASE 6 INHIBITOR	CELL CYCLE INHIBITOR P18-
Itiph A 18 164 5.40E-25 0.31 0.93 CYCLIN-DEPENDENT										CHAIN: A, B;	INHIBITOR, P18-INK4C(INK6),
Litho A 18 164 5.40E-25 0.31 0.93 CYCLIN-DEPENDENT											ANKYRIN REPEAT, 2 CDK 4/6 INHIRITOR
Likh A 75 232 5.40E-36 63.76 CYCLIN-DEPENDENT Likh A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Likh D 107 238 3.60E-26 -0.33 0.33 CYCLIN-DEPENDENT Likh D 107 238 3.60E-36 -0.33 0.33 CYCLIN-DEPENDENT Likh D 107 238 3.60E-36 -0.33 0.33 CYCLIN-DEPENDENT Likh D 13 177 3.60E-33 -0.04 0.9 CYCLIN-APA-B-ALPHA; CHAIN: C; I Likh D 21 216 7.20E-39 CHAIN: A; INF-KAPPA-B P65 SUBUNIT; CHAIN: C; I Likh D 39 228 7.20E-39 COS Likh D 5 144 1.30E-27 -0.21 0.76 CHAIN: A; INF-KAPPA-B P65 SUBUNIT; CHAIN: C; I KAPPA-B-ALPHA; CHAIN: C; I KAPPA-		lihb	4	18	164	5.40E-25	0.31	0.93		CYCLIN-DEPENDENT	CELL CYCLE INHIBITOR P18-
Litho A 75 232 5.40E-36 G3.76 CYCLIN-DEPENDENT Litho A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Litho A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Litho D 107 238 3.60E-26 -0.33 0.33 OHAIN: A. BrKAPPA-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. P. CAPAB-B P65 SUBUNIT; CHAIN: CI. CHAIN: A. BrKAPPA-B P65 SUBUNIT; CHAIN: CI. CHAIN: A. BrKAPPA-B P65 SUBUNIT; CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CAPBA-B P65 SUBUNIT; CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CAPBA-B P65 SUBUNIT; CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CAPBA-B P65 SUBUNIT; CHAIN: CI. CHAIN: CI. CHAIN: CI. CHAIN: CI. CAPBA-B P65 SUBUNIT; CHAIN: CI. CHAIN: CHAIN: CI. CHAIN: CHAIN: CI. CHAIN: CHA										KINASE 6 INHIBITOR;	INK4C(INK6); CELL CYCLE
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Liph A 75 232 5.40E-36 63.76 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT Liph A 78 232 5.40E-36 0.33 0.33 O.44IN: A, INF-KAPPA-B-P50 Liph A 78 238 3.60E-26 -0.33 0.33 O.44IN: C, I Liph A 78 238 3.60E-33 -0.04 0.9 O.44IN: A, INF-KAPPA-B-P50 Liph A 78 228 7.20E-39 0.05 O.69 O.44IN: A, INF-KAPPA-B-P50 Liph A 78 228 7.20E-39 0.05 O.69 O.44IN: A, INF-KAPPA-B-P50 Liph A 78 228 7.20E-39 O.05 O.69 O.44IN: A, INF-KAPPA-B-P50 Liph A 78 144 1.30E-27 -0.21 0.76 O.76											ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR
Iikh A 78 232 5.40E-36 0.24 1 CYCLIN-DEPENDENT CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: B, B; C		lihb	A	75	232	5.40E-36			63.76	CYCLIN-DEPENDENT	CELL CYCLE INHIBITOR P18-
Iikn D 107 238 3.60E-36 0.24 1 CYCLIN-DEPENDENT Iikn D 107 238 3.60E-36 -0.33 0.33 NF-KAPPA-B P6S SUBUNIT; CHAIN: C, I.P. KAPPA-B P50D SUBUNIT; CHAIN: C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C, I.P. KAPPA-B P6S SUBUNIT; C,										KINASE 6 INHIBITOR;	NK4C(NK6); CELL CYCLE
Ilikn D 39 228 7.20E-39 0.24 1 CYCLIN-DEPENDENT										CHAIN: A, B;	INHIBITOK, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6
Likn D 107 238 3.60E-26 -0.33 0.33 OF-CLIN-DEPENDENT CHAIN: A, B; CHAIN: A, B; CHAIN: A, B; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I - KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I - KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I - KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I - KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I - KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I - KAPPA-B-ALPHA; CHAIN: C, I - CANDA-C, C											INHIBITOR
IIKn D 107 238 3.60E-26 -0.33 0.33 O.4ADPA-B P65 SUBUNIT;		lihb	۷	18	232	5.40E-36	0.24	_		CYCLIN-DEPENDENT	CELL CYCLE INHIBITOR P18-
IIKn D 107 238 3.60E-26 -0.33 0.33 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I-										KINASE 6 INHIBITOR;	INK4C(INK6); CELL CYCLE
likn D 107 238 3.60E-26 -0.33 0.33 NF-KAPPA-B P65 SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B-ALPHA; CHAIN: D; I-KAPPA-B P65 SUBUNIT; CHAIN: C; I-KAPPA-B P65 SUBUNIT; CH		_								CHAIN: A, B;	INHIBITOR, P18-INK4C(INK6),
likn D 107 238 3.60E-26 -0.33 0.33 NF-KAPPA-B P65 SUBUNIT; likn D 13 177 3.60E-33 -0.04 0.9 NF-KAPPA-B P65 SUBUNIT; likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 NF-MA											ANKYRIN REPEAT, 2 CDK 4/6
13 177 3.60E-33 -0.04 0.9 NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P65 SUBUN		Likn		107	238	3 60F-26	-0 33	0 33		NE-KAPPA, B P64 SUBUNIT.	TP ANSCRIPTION EACTOR PASS DAND
Iikn D 13 177 3.60E-33 -0.04 0.9 NF-KAPPA-B-ALPHA, CHAIN: D;	_				<u> </u>	1	}			CHAIN: A: NF-KAPPA-B P50D	TRANSCRIPTION FACTOR, IKB/NFKB
likn D 13 177 3.60E-33 -0.04 0.9 NF-KAPPA-B-R6S SUBUNIT; likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: C; I- KAPPA-B; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B P65 SUBUNIT;	_									SUBUNIT; CHAIN: C; I-	COMPLEX
likn D 13 177 3.60E-33 -0.04 0.9 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: C; I- KAPPA-B-A; NF-KAPPA-B P50D CHAIN: A; NF-KAPPA-B P50D CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B P65 SUBUNIT;	\neg									KAPPA-B-ALPHA; CHAIN: D;	
likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P50D likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT;			Ω	13	177	3.60E-33	-0.04	6.0		NF-KAPPA-B P65 SUBUNIT;	TRANSCRIPTION FACTOR P65; P50D;
likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D										CHAIN: A; NF-KAPPA-B P50D	TRANSCRIPTION FACTOR, IKB/NFKB
likn D 21 216 7.20E-39 69.17 NF-KAPPA-B P65 SUBUNIT; CHAIN: A: NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B P65 SUBUNIT;										KAPPA-B-ALPHA: CHAIN: D:	COMPLEX
CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; I- KAPPA-B-ALPHA; CHAIN: D; I- KAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT: CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; I- KAPPA-B-ALPHA; CHAIN: D; I- KAPPA-B-B65 SUBUNIT; CHAIN: C; I- KAPPA-B-P65 SUBUNIT; CHAIN: C; I- KAPPA-B-P65 SUBUNIT; CHAIN: C; I- KAPPA-B-P65 SUBUNIT; CHAIN: C; I- KAPPA-B-P65 SUBUNIT; CHAIN: C; I- KAPPA-B-P65 SUBUNIT; CHAIN: C; I- KAPPA-B-P50D SUBUNIT; CHAIN: C; I	_	likn	Q	21	216	7.20E-39			69.17	NF-KAPPA-B P65 SUBUNIT;	TRANSCRIPTION FACTOR P65; P50D;
likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B-ALPHA, CHAIN: D; likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P50D SUBUNIT; CHAIN: C, I- KAPPA-B P50D SUBUNIT; CHAIN: C, I- CHAIN: C, I- CHAIN: C, I-										CHAIN: A; NF-KAPPA-B P50D	TRANSCRIPTION FACTOR, IKB/NFKB
likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P65 SUBUNIT; Iikn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT;	_				_					SUBUNIT; CHAIN: C; I-	COMPLEX
likn D 39 228 7.20E-39 -0.05 0.69 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT: CHAIN: C; I-KAPPA-B P60D KAPPA-B-ALPHA; CHAIN: D; likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B P50D	7									KAPPA-B-ALPHA, CHAIN: D,	
CHAIN: A; NF-KAPPA-B P50D CHAIN: A; NF-KAPPA-B P50D SUBUNIT: CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I-	_	ik i	Ω	39	228	7.20E-39	-0.05	69.0		NF-KAPPA-B P65 SUBUNIT,	TRANSCRIPTION FACTOR P65, P50D;
Iikn D 5 144 1.30E-27 -0.21 0.76 RAPPA-B-ALPHA; CHAIN: D; CHAIN: A; NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D										CHAIN: A; NF-KAPPA-B P50D	TRANSCRIPTION FACTOR, IKB/NFKB
likn D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B-ALPHA, CHAIN: D; CHAIN: A; NF-KAPPA-B P50D CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D	_									SUBUNIT: CHAIN: C; I-	COMPLEX
18h D 5 144 1.30E-27 -0.21 0.76 NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CAPAN: C. 1-	7	1								KAPPA-B-ALPHA; CHAIN: D;	
		lika L	Ω	S	44	1.30E-27	-0.21	0.76		NF-KAPPA-B P65 SUBUNIT,	TRANSCRIPTION FACTOR P65, P50D,
_	_									CHAIN: A; NF-KAPPA-B P50D	TRANSCRIPTION FACTOR, IKB/NFKB
										SUBUNII; CHAIN: C; I-	COMPLEX

PDB annotation	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK-REPEAT	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK-REPEAT	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK	REPEAT), ANKYRIN Z KEFEAT HELIX COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK)	KEFEAT), ANKTRING TREFAT HELLA COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT) ANKYRN 2 REPEAT HELLX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (ANTI- ONCOGENE/ANKYRIN REPEATS) P53BP2: ANKYRIN REPEATS, SH3, P53, TUMOR SUPPRESSOR, MULTIGENE 2 FAMILY, NUCLEAR PROTEIN, PHOSPHORYLATION, DISEASE MUTATION, 3 POLYMORPHISM, COMPLEX (ANTI- ONCOGENE/ANKYRIN REPEATS)		COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI: COMPLEX
Coumpound	MYOTROPHIN; CHAIN: NULL	MYOTROPHIN; CHAIN: NULL	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA;	CHAIN: E, F; NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA;	CHAIN: E. F. NF-KAPPA-B P65; CHAIN: A. C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: F. E.	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; LKAPPA-B-ALPHA; CHAIN: F.	P53; CHAIN: A; 53BP2; CHAIN: B;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1:
SeqFold score		64.15				65.66	64.29		
PMF score	_		0.94	0.92	0.78			-	_
Verify score	0.14		0.19	-0.07	0.07			0.79	0.83
PSI- BLAST	2.60E-28	6.50E-33	9.00E-26	9.00E-33	1.10E-38	1.10E-38	1.60E-20	7.80E-42	2.60E-42
End	225	192	238	177	228	202	239	312	147
Start	110	75	901	12	39	7	74	191	5.
Chain ID			EL	ш	E	ш	В	B	В
PDB ID	1my	1my o	Infi	Infi	- Infi	lnfi	lycs	lawc	lawc
SEQ ID	719	719	719	719	719	719	719	721	721

SEQ TD NO:	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
									CHAIN: B; DNA; CHAIN: D, E;	(TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
721	lawc	B _	2	147	3.60E-34	0.71	_		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
721	lawc	В	226	378	5.20E-43	1.14	_		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
721	lawc	B	23	180	1.30E-44	0.72	_		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
721	lawc	В	231	378	3.60E-39	0.9	-		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
721	lawc	8	259	412	5.20E-45	1.23	_		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION

PDB annotation	REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, NHY YRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI: COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2
	REGULATI NUCLEAR ANKYRIN 3 FACTOR	COMPLEX REGULATI GABPBET/ (TRANSCR REGULATI NUCLEAR ANKYRIN	COMPLEX REGULATI GABPBETA (TRANSCR REGULATI NUCLEAR ANKYRIN 3 FACTOR	COMPLEX REGULATI GABPBET/ (TRANSCR REGULATI NUCLEAR ANKYRIN 3 FACTOR	COMPLEX REGULAT GABPBET, (TRANSCR REGULAT NUCLEAR ANKYRIN 3 FACTOR	COMI REGL GABF (TRA)
Coumpound		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A: GA BINDING PROTEIN BETA I; CHAIN: B: DNA; CHAIN: D. E;	GA BINDING PROTEIN ALPHA: CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAM: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;
SeqFold score						
PMF		-	-		-	0.62
Verify score		0.93	0.93	0.69	0.67	0.22
PSI- BLAST		5.40E-41	1.80E-36	1.30E-38	1.80E-36	7.20E-32
End AA		411	442	180	475	487
Start		264	297	33	330	363
Chain ID		В	m	B	B	æ
PDB ID		lawc	lawc	lawc	lawc	lawc
SEQ ID NO:		721	721	721	721	721

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify score	PMF	SeqFold score	Coumpound	PDB annotation
.1										NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
	lawc	æ	61	213	6.50E.47	0.59	1		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
	lawc	B	99	213	1.40E-39	0.79	-		GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR
721	lawc	В	66	246	5.40E-38	0.87			GA BINDING PROTEIN ALPHA: CHAIN: A; GA BINDING PROTEIN BETA I; CHAIN: B; DNA; CHAIN: D, E;	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR
	1bd8		128	281	2.60E-38	6.79	-		P19NK4D CDK4/6 INHIBITOR; CHAIN: NULL;	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF
721	1 bd8		191	348	2.60E-38	0.12	0.42		P19TNK4D CDK4/6 INHIBITOR; CHAIN: NULL;	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF
721	15d8			149	1.20E-40	0.43	-		P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF
	1bd8		727	381	9.10E-41	0.73	1		P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF
721	1bd8		24	182	1.30E-41	0.53	_		P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR,

PDB Chain ID ID		Start	End AA	PSI- BLAST	Verify score	PMF score	SeqFold	Coumpound	PDB annotation
									ANKYRIN MOTIF
263	263		414	1.20E-38	0.89	_		P19NK4D CDK4/6 INHIBITOR; CHAIN: NULL;	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF
297	297	}	442	1.40E-30	0.74	_		P19INK4D CDK4/6	TUMOR SUPPRESSOR TUMOR
								INHIBITOR; CHAIN: NULL;	SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF
62	62		215	7.80E-44	19.0	_		P19INK4D CDK4/6	TUMOR SUPPRESSOR TUMOR
								INHIBITOR; CHAIN: NULL;	SUPPRESSOR, CDK4/6 INHIBITOR, ANK YRIN MOTIF
B 132	132		285	1.30E-39	0.83	_		CYCLIN-DEPENDENT	COMPLEX (INHIBITOR
								KINASE 6; CHAIN: A;	PROTEIN/KINASE) INHIBITOR
_								P19INK4D; CHAIN: B;	PROTEIN, CYCLIN-DEPENDENT
									ALPHA/BETA, COMPLEX (INHIBITOR
									PROTEIN/KINASE)
B 163	163		350	5.20E-38	0.13	_	-1-	CYCLIN-DEPENDENT	COMPLEX (INHIBITOR
								KINASE 6; CHAIN: A;	PROTEIN/KINASE) INHIBITOR
								P19INK4D; CHAIN: B;	PROTEIN, CYCLIN-DEPENDENT
					_				NINASE, CELL CYCLE Z CONTROL,
									ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
B 2	2		153	1.00E-41	0.81	_		CYCLIN-DEPENDENT	COMPLEX (INHIBITOR
								KINASE 6; CHAIN: A;	PROTEINKINASE) INHIBITOR
								P19INK4D; CHAIN: B;	PROTEIN, CYCLIN-DEPENDENT
									KINASE, CELL CYCLE 2 CONTROL,
									ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)
В 22	22		185	6.50E-43	0.61	_		CYCLIN-DEPENDENT	COMPLEX (INHIBITOR
								KINASE 6; CHAIN: A;	PROTEIN/KINASE) INHIBITOR
,								P19INK4D; CHAIN: B;	PROTEIN, CYCLIN-DEPENDENT
									KINASE, CELL CYCLE 2 CONTROL,
_						_			ALPHA/BETA, COMPLEX (INHIBITOR
	555	Ī	200	., 200					PROTEIN/KINASE)
067	067		385	1.30E-43	0.78	_		CYCLIN-DEPENDENT	COMPLEX (INHIBITOR
						_		KINASE 6; CHAIN: A;	PROTEIN/KINASE) INHIBITOR
		_						PI9INK4D; CHAIN: B;	PROTEIN, CYCLIN-DEPENDENT
									KINASE, CELL CYCLE 2 CONTROL,
									ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/VINASE)
B 263	263	-	416	5.20E-41	1.13			CYCLIN-DEPENDENT	COMPLEX (NHIBITOR
	1			1				מונים מונים מונים מונים מונים	יייייייייייייייייייייייייייייייייייייי

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PDB annotation	PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/CROWTH FACTOR	HORMONE/GROWTH FACTOR P18-INK4C; CELL CYCLE INHIBITOR, P18NK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT	SIGNALING PROTEIN HELIX-TURN- HELIX, ANKYRIN REPEAT
Coumpound	KINASE 6; CHAIN: A; PI9INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A:	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;	CYCLIN-DEPENDENT KINASE 4 INHIBITOR B; CHAIN: A;
SeqFold score					90.26					
PMF		_	-	0.35		_	-	_	-	
Verify		0.39	0.61	0.37		9.0	0.66	0.58	0.65	0.61
PSI- BLAST		6.50E-43	7.20E-38	1.80E-32	7.20E-38	6.50E-36	1.30E-37	1.30E-40	2.60E-38	2.60E-38
End		252	185	480	258	153	384	416	185	216
Start AA		64	33	330	16	14	250	283	52	84
Chain ID		В	4	∢	<	∢	V	∢	∢	∢
PDB ID		1blx	1bu9	1bu9	15u9	149s	s6p1	149s	149s	149s
SEQ NO:		721	721	721	721	721	721	721	721	721

d PDB annotation	NT CELL CYCLE INHIBITOR P18- NK, INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR				1	JEGNATOR TRANSCRIPTION FACTOR P65; P50D; A-B P50D TRANSCRIPTION FACTOR, IKBNIFKB COMPLEX J. I. COMPLEX	ABUNIT; TRANSCRIPTION FACTOR P65; P50D; AB P50D TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX THAIN. D.	JBUNIT; TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX COMPLEX	BUNIT; TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX
ld Coumpound	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-AI PHA: CHAIN: D:	NP-KAPPA-B F65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B AI PHA: CUAN: D.	NF-KAPPA-B F65 SUBUNIT; CHAIN: 4, NF-KAPPA-B P50D SUBUNIT; CHAIN: C; 1- KAPPA-B-AI, PHA: CHAIN: D:	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-AI PHA: CHAIN: D:	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; 1-
F SeqFold				88.22	82.11				
PMF	-	_	0.92	_		0.93	0.99	-	
Verify score	0.7	0.55	0.5			-0.02	0.48	0.56	0.55
PSI- BLAST	1.80E-32	3.60E-37	1.30E-31	3.60E-37	2.60E-57	2.60E-55	5.20E-52	5.40E-43	7.80E-52
End	446	184	479	250	333	386	188	411	416
Start	297	33	330	96	127	191	2	226	226
Chain ID	<	∢	4	∢	Ω	Ω	Q	Ω	D
PDB ID	1 ihb	1ihb	lihb	libb	likn	1 ikn	likn	i E	l ikn
SEQ ID NO:	721	721	721	721	721	721	121	721	127

PDB annotation	TRANSCRIPTION FACTOR, IKBNFKB COMPLEX	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX		TRANSCRIPTION FACTOR P65, P50D, TRANSCRIPTION FACTOR, IKBANFKB COMPLEX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANK YRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANK YRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANK YRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION
Coumpound	CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; L-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF.KAPPA-B P65, CHAIN: A, C; NF.KAPPA-B P50, CHAIN: B, D; I.KAPPA-B-ALPHA; CHAIN: E, F;	NF.KAPPA-B P65; CHAIN: A, C; NF.KAPPA-B P50; CHAIN: B, D; 1-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65; CHAIN: A,
SeqFold score					85.42						
PMF score		0.05	-	-		-		-	-	_	_
Verify		0.09	0.2	0.22		0.08	69.0	0.49	0.82	8.0	0.65
PSI- BLAST		7.20E-34	5.40E-38	2.60E-57	2.60E-52	2.60E-52	7.80E-42	7.80E-53	7.20E-43	6.50E-52	1.60E-38
End		486	233	256	322	350	153	216	411	420	475
Start		325	19	19	124	124	2	21	224	226	292
Chain ID		Q	Q	Q	ங	ய	ш	m	ш	ជា	ы
PDB ID		likn	Likn	l ikn	Infi	Infi	1nfi	Infi	Infi	Infi	Infi
SEQ NO:		721	721	721	721	721	721	721	721	721	721

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PDB annotation	(TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX	CHARLOS TOOM ON BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA I: COMPLEX	(TRANSCRIPTION) DNA-BINDING, 2	NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION	3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA;	GABPBETA1; COMPLEX (TRANSCRIPTION	REGULATION/DNA), DNA-BINDING, 2	ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGILI ATION MAN GARPAI PHA	GABPBETAI; COMPLEX	(IKANSCKIPIJON REGIII ATTON/DNA) DNA-BINDING 2	NUCLEAR PROTEIN, ETS DOMAIN.	ANKYRIN REPEATS, TRANSCRIPTION	COMPLEX (TRANSCRIPTION
Coumpound	B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65, CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B: DNA: CHAIN: D. E.		GA BINDING PROTEIN ALPHA: CHAIN: A; GA BINDING PROTEIN BETA 1:	CHAIN: B; DNA; CHAIN: D, E;			GA BINDING PROTEIN ALPHA; CHAIN: A; GA	BINDING PROTEIN BETA 1; CHAIN: B: DNA: CHAIN: D. E:			GA BINDING PROTEIN AT PHA: CHAIN: A: GA	BINDING PROTEIN BETA 1;	CHAIN: B; DNA; CHAIN: D, E;			GA BINDING PROTEIN
SeqFold score													64.41			_		
PMF		96.0	_		0.64				-0.17					_				_
Verify score		0.52	0.32		0.4				0.02									0.35
PSI- BLAST		1.80E-33	1.60E-31		1.60E-23				3.60E-30				5.40E-36					5.40E-36
End AA		486	280		343				408				198					157
Start AA		323	107		175				265				31					9
Chain ID		ங	В		В				<u>m</u>				В					В
PDB ID		lnfi	lawc		lawc				lawc				lawc					lawc
SEQ ID NO:		721	723		723				723				723					723

PDB annotation	GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	TUMOR SUPPRESSOR TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE NHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH FACTOR P18-
Coumpound	BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	PI9INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A:	CYCLIN-DEPENDENT
SeqFold score			56.93						
PMF		-		1	1	,	_	0.3	96.0
Verify score		0.71		0.43	0.21	0.44	0.44	0.24	0.25
PSI- BLAST		2.60E-27	3.60E-29	7.20E-27	3.60E-29	2.60E-26	5.40E-29	3.60E-29	5.40E-34
End		212	201	182	160	202	148	291	162
Start AA		72	38	43	6	43	6	107	9
Chain ID		В				В	В	¥	V
PDB ID		lawc	1 pq8	8pq1	8pq1	1blx	1bix	1bu9	1bu9
SEQ ID NO:		723	723	723	723	723	723	723	723

	PDB ID	Chain ID	Start AA	End	PSI- BLAST	Verify score	PMF score	SeqFold score	Coumpound	PDB annotation
-	+								KINASE 6 INHIBITOR; CHAIN: A;	NK4C; CELL CYCLE NHIBITOR, P18NK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONEGROWTH FACTOR
1 bu9		Y	73	234	1.80E-26	0.32	0.88		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR
dii.	<u> </u>	V	107	284	1.80E-28	0.07	0.35		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR
<u>-</u>	Lihb	4	9	191	1.80E-33	0.15	0.99		CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CELL CYCLE INHIBITOR P18- INK4C(INK6); CELL CYCLE INHIBITOR, P18-INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR
lik Ik		D	2	174	1.80E-43	0.13	_		NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-AI, PHA: CHAIN: D;	TRANSCRIPTION FACTOR P65, P50D: TRANSCRIPTION FACTOR, IKBNFKB COMPLEX
32		D	2	208	1.80E-43			61.01	NF-KAPPA-B P65 SUBUNIT: CHAIN: A: NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKBNFKB COMPLEX
5 1		Ω	35	229	3.60E-36	0.37	0.95		NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; I- KAPPA-B-ALPHA; CHAIN: D;	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX
ik k	_	Ω	89	280	1.10E-28	-0.02	0.43		NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF-KAPPA-B P50D SUBUNIT; CHAIN: C; J- KAPPA-B-ALPHA; CHAIN: D;	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX
Infi		យ	136	288	5.40E-30	90.0	0.65		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: F. F.	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT) ANX YEN 2 REPEAT HE! Y
ا بسا	Infi	ய	2	174	9.00E-44	0.31	-		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN:	COMPLEX (TRANSCRIPTION REGANK REPEAT) COMPLEX

PDB annotation	(TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANK YRIN 2 REPEAT HELIX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITORNUCLEASE) COMPLEX (INHIBITORNUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS
Coumpound	B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	NF-KAPPA-B P65, CHAIN: A, C, NF-KAPPA-B P50, CHAIN: B, D, I-KAPPA-B-ALPHA; CHAIN: E, F:	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B-ALPHA; CHAIN: E, F;	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A. D: ANGIOGENIN; CHAIN: B, E:	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;
SeqFold score			59.74				93.46	
PMF				0.07	0.16	0.55		_
Verify score		0.49		-0.4	-0.3	-0.33		0.15
PSI- BLAST		5.40E-36	9.00E-44	1.30E-13	9.00E-17	2.60E-22	3.90E-36	9.10E-39
End AA		229	246	369	577	597	558	376
Start		34	36	-	219	267	82	83
Chain ID		ഖ	ы	∢	∢	¥	V	∢
PDB ID		Infi	Infi	la4y	la4y	la4y	la4y	1a4y
SEQ ID NO:		723	723	725	725	725	725	725

SEQ ID	PDB ID	Chain ID	Start	End	PSI- BLAST	Verify	PMF	SeqFold score	Coumpound	PDB annotation
ÖZ			:)65	1000	20			TIS DATA HATDRIN IN. CHARL.	COMPLEY ALLOT EAD DEOTEIN BNA)
725	la9n	∢	134	306	5.20E-27	70.02	0.57		02 KNA HAIKFIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 p": CUAN: B. D:	COMPLEX (NOCLEAR PROTEIN/RIA) COMPLEX (NUCLEAR PROTEIN/RIA), PNA SNRNP PIRONICI FOPROTFIN
77.5	1,000	<	203	351	1 30E 25	90.0	0.75		112 PNA HAIRPIN IV: CHAIN:	COMPLEX (NICLEAR PROTEIN/RNA)
3	14711	<	607		1.300-1	3	3.5		O. R. U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	1a9n	¥	430	576	3.90E-19	0.12	0.77		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									0, R; U2 A; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	¥	454	576	6.50E-20	-0.33	0.45		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	٧	68	228	3.90E-29	0.22	0.99		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
			_						Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	ပ	134	618	1.20E-27	-0.06	0.84		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	ပ	203	351	2.60E-25	0.24	0.88		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	ပ	430	576	3.90E-19	-0.04	69.0		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									Q, R; U2 A; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	ပ	454	576	6.50E-20	0	0.45		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	ပ	82	213	9.10E-26	0.05	0.55		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
			-						Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	la9n	ပ	68	236	6.50E-29	0.27	96.0		U2 RNA HAIRPIN IV; CHAIN:	COMPLEX (NUCLEAR PROTEIN/RNA)
									Q, R; U2 A'; CHAIN: A, C; U2	COMPLEX (NUCLEAR PROTEIN/RNA),
									B"; CHAIN: B, D;	RNA, SNRNP, RIBONUCLEOPROTEIN
725	140p	4		158	1.60E-26	-0.09	89.0		INTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE RICH
_				_						REPEAT, CALCIUM BINDING, CELL
										ADHESION
725	1406	¥	104	319	1.30E-24	0.02	_		INTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE RICH
										REPEAT, CALCIUM BINDING, CELL ADHESION
725	140b	¥	216	364	5.40E-29	0.12	_		INTERNALIN B; CHAIN: A;	CELL ADHESION LEUCINE RICH REPFAT CALCII M RINDING CFI I
										מממט לטיוום וווסוסמטילי ועד וחו

PDB annotation	ADHESION	A; CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION		A; CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION		ANSF FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT	-	ANSF STRUCTURE, RAB F:		ANSF FORMYLMETHIONINE, ALPHA		┢	ANSF STRUCTURE, RAB					ANSF STRUCTURE, RAB F. GERANYI GERANYI TRANSFERASE		ANSF FORMYLMETHIONINE, ALPHA	SUBUNIT, BETA SUBUNIT	TRANSFERASE CRYSTAL	_
Coumpound		INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	RAB GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT; CHAIN: A, C; RAB	GERANYLGERANYLTRANSF ERASE BETA SUBUNIT; CHAIN: B, D;	RAB	GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT:	CHAIN: A. C; RAB	GERANYLGERANYLTRANSF	CHAIN: B, D;	RAB	GERANYLGERANYLTRANSF	CHAIN: A. C. RAB	GERANYLGERANYLTRANSF	CHAIN: B, D;	RAB	GERANYLGERANYLTRANSF	CHAIN: A. C. RAB	GERANYLGERANYLTRANSF	ERASE BETA SUBUNIT;	RAB	GERANYLGERANYLTRANSF
SeqFold score			:																				
PMF		0.4	0.92	0.34	_		0.95					96.0					0.22					0.07	
Verify score		0.04	0.18	60:0	0.12		-0.23					0.18					0.08					0.07	
PSI- BLAST		9.00E-28	5.40E-24	1.10E-21	3.60E-13		5.40E-13					9.00E-13	· · · · · ·				1.80E-13					7.20E-09	
End		406	166	965	301		346					369					533					109	
Start		242	32	415	216		246					268			_		428					475	
Chain ID		4	4	V	V		A					4					Α					\ \ \	
PDB D		1406	1406	1006	1dce		1dce					1dce					1dce					1dce	
SEQ ID	Ö	725	725	725	725		725					725					725					725	

PDB annotation	2.0 A 2 RESOLUTION, N- FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT	TRANSFERASE CRYSTAL STRUCTURE, RAB GERANYLGERANYLTRANSFERASE, 2.0 A 2 RESOLUTION, N- FORMYLMETHIONINE, ALPHA SUBUNIT, BETA SUBUNIT	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE. RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	CONTRACTILE PROTEIN LEUCINE- RICH REPEAT, BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FLAGELLA	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN
	2.0 A 2 RESOLUTION, N. FORMYLMETHIONINE, SUBUNIT, BETA SUBUN	TRANSFERASE CR STRUCTURE, RAB GERANYLGERAN 2.0 A 2 RESOLUTIC FORMYLMETHION SUBUNIT, BETA SI	CONTRACTILE PROT RICH REPEAT, BETA-CYLINDER, DYNEIN, CHLAMYDOMONAS,	CONTRACTILE PROTEI RICH REPEAT, BETA-B CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, F	CONTRACTILE PROTE RICH REPEAT, BETA-B CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, F	CONTRACTILE PROTEI RICH REPEAT, BETA-B. CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, FI	CONTRACTILE PROTE RICH REPEAT, BETA-B CYLINDER, DYNEIN, 2 CHLAMYDOMONAS, F	LIGASE CY ASSOCIATE A/CDK2-AS; SKP1, SKP2, RICH REPE, UBIQUITIN	ASSOCIATE
Coumpound	CHAIN: A, C; RAB GERANYLGERANYLTRANSF ERASE BETA SUBUNIT; CHAIN: B, D:	RAB GERANYLGERANYLTRANSF ERASE ALPHA SUBUNIT; CHAIN: A, C; RAB GERANYLGERANYLTRANSF ERASE BETA SUBUNIT; CHAIN: B, D;	OUTER ARM DYNEIN; CHAIN: A;	OUTER ARM DYNEIN: CHAIN: A;	OUTER ARM DYNEIN; CHAIN: A;	OUTER ARM DYNEIN; CHAIN: A;	OUTER ARM DYNEIN; CHAIN: A;	SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	SKP2; CHAIN: A, C, E, G, 1, K, M, O; SKP1; CHAIN: B, D, F,
SeqFold score									
PMF score		0.95	0.21	0.94	0.12	0.52	0.75	0.04	0.09
Verify		0.3	-0.34	-0.36	-0.73	-0.47	-0.79	0.02	-0.35
PSI- BLAST		1.30E-13	3.60E-13	1.10E-14	7.20E-13	1.30E-13	3.90E-26	3.60E-07	1.30E-09
End		170	140	324	346	164	213	172	575
Start		09	17	210	232	70	82	_	264
Chain ID			V	V	V	V	V V	∢	∀
PDB ID		1dce	6sp1	1489	1ds9	1ds9	6sp1	1 fqv	1 fqv
SEQ ID	O	725	725	725	725	725	725	725	725

PDB annotation	RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, LRR, LEUCINE- RICH REPEAT, SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45, CYCLIN A/CDK2- ASSOCIATED P19, SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	LIGASE CYCLIN A/CDK2- ASSOCIATED P45; CYCLIN A/CDK2- ASSOCIATED P19; SKP1, SKP2, F-BOX, LRRS, LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	TRANSCRIPTION RNAIP; RANGAP; GTPASE-ACTIVATING PROTEIN FOR SPII, GTPASE-ACTIVATING PROTEIN, GAP, RNAIP, RANGAP, LRR, LEUCINE-2 RICH REPEAT PROTEIN, TWINNING, HEMIHEDRAL TWINNING, 3 MEROHEDRAL TWINNING, MEROHEDRAL	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE-
Coumpound		SKP2; CHAIN: A, C, E, G, I, K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	SKP2; CHAIN: A, C; SKP1; CHAIN: B, D;	GTPASE-ACTIVATING PROTEIN RNA1_SCHPO; CHAIN: A, B;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;
SeqFold								
PMF score		0.06	0.11	0.27	0.31	0.04	0.22	0.87
Verify score		-0.06	-0.09	-0.12	-0.14	-0.22	-0.07	-0.26
PSI- BLAST		1.00E-16	3.60E-11	7.80E-19	6.50E-40	1.30E-09	1.10E-19	5.40E-21
End		253	453	577	338	431	498	592
Start		88	244	401	68	211	110	239
Chain ID		<	⋖	<	∢	∢		
PDB ID		1 fqv	1652	1652	1fs2	lyrg	2bnh	2bnh
SEQ ID NO:		725	725	725	725	725	725	725

	FOR, UCINE-	TOR, UCINE-	HA; DING, 2 AIN, LPTION	HA; DING, 2 AIN, SIPTION	HA; DING, 2 IAIN, SIPTION	HA; DING, 2
notation	RICH REPEATS ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGENIN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA, DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI: COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BINDING, 2
PDB annotation	EATS ATION RNA CLEASE/AN OR ACETYI	ATION RNATION	COMPLEX (TRANSCRIPTION REGULATIONDNA) GABPAL GABPBETAI; COMPLEX (TRANSCRIPTION REGULATIONDNA), DINA-BI NUCLEAR PROTEIN, ETS DO ANK YRIN REPEATS, TRANS(3) FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPAL GABPBETA1; COMPLEX TRANSCRIPTION REGULATION/DNA), DNA-BI NUCLEAR PROTEIN, ETS DO ANKYRIN REPEATS. TRANS 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPAL GABPBETA!: COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA-BI NUCLEAR PROTEIN, ETS DO ANK YRIN REPEATS, TRANSG 5 FACTOR	COMPLEX (TRANSCRIPTION REGULATIONIDNA) GABPAL GABPBETA1; COMPLEX (TRANSCRIPTION REGULATIONIDNA), DINA-BI
	RICH REPEATS ACETYLATION RIBONUCLEASI INHIBITOR ACE	ACETYLATION RIBONUCLEASI INHIBITOR ACE RICH REPEATS	COMPLEX REGULATI GABPBET/ (TRANSCR REGULATI NUCLEAR ANKYRIN 3 FACTOR	COMPLEX REGULATI GABPBET ((TRANSCR REGULATI NUCLEAR ANK YRIN 3 FACTOR	COMPLEX REGULATI GABPBET/ (TRANSCR REGULATI NUCLEAR ANKYRIN 3 FACTOR	COMPLE REGULA GABPBE (TRANS(REGULA
	IIBITOR:	IIBITOR;	IN 3A 3ETA 1; AIN: D, E;	IIN 3A 3ETA I; AIN: D, E;	SIN SA BETA 1; AIN: D, E;	SIN SA BETA 1; AIN: D, E;
Coumpound	RIBONUCLEASE INHIBITOR; CHAIN: NULL;	RIBONUCLEASE INHIBITOR; CHAIN: NULL;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA, CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;
0	RIBONUCLEAS CHAIN: NULL;	RIBONUCLEAS CHAIN: NULL;	GA BINDI ALPHA; C BINDING CHAIN: B;	GA BINDI ALPHA; C BINDING CHAIN: B	GA BINDI ALPHA; C BINDING CHAIN: B	GA BINDI ALPHA, C BINDING CHAIN: B
SeqFold score		90.38			53.5	
PMF score	10.0	6	-0.03	0.29		0.72
Verify	-0.33		0.12	0.19		0.32
PSI. BLAST	1.80E-19	1.10E-19	3.60E-32	1.30E-35	1.30E-35	1.60E-31
End	448	558	145	167	179	203
Start	35	82	2	23	27	\$\$
Chain ID			В	В	В	B
PDB ID	2bnh	2bnh	lawc	lawc	lawc	lawc
SEQ D NO.	725	725	728	728	728	728

SEQ ID	PDB TD	Chain ID	Start	End	PSI- BLAST	Verify score	PMF score	SeqFold	Coumpound	PDB annotation
ÖZ										3 FACTOR
728	1bu9	<	23	183	1.80E-29	0.33	0.01		CYCLIN-DEPENDENT KINASE 6 INHIBITOR;	HORMONE/GROWTH FACTOR P18- INK4C; CELL CYCLE INHIBITOR,
								_	CHAIN: A;	P18INK4C, TUMOR, SUPPRESSOR.
										HORMONE/GROWTH FACTOR
728	1bu9	4	5	150	1.80E-29	0.28	60.0		CYCLIN-DEPENDENT	HORMONE/GROWTH FACTOR P18-
									KINASE 6 INHIBITOR;	INK4C; CELL CYCLE INHIBITOR,
									CHAIN: A;	P18INK4C, TUMOR, SUPPRESSOR,
										CICLIN- 2 DEFENDENT NINASE, HORMONE/GROWTH FACTOR
728	1 bu9	<	97	250	1.40E-27	0.04	-0.15		CYCLIN-DEPENDENT	HORMONE/GROWTH FACTOR P18-
									KINASE 6 INHIBITOR;	NK4C; CELL CYCLE INHIBITOR,
									Crain: A;	CYCLIN-2 DEPENDENT KINASE.
										HORMONE/GROWTH FACTOR
728	1 ihb	4	23	991	3.60E-29	0.25	0.41		CYCLIN-DEPENDENT	CELL CYCLE INHIBITOR P18-
					-				KINASE 6 INHIBITOR;	INK4C(INK6); CELL CYCLE
									CHAIN: A, B;	INHIBITOR, P18-INK4C(INK6),
										ANKIKIN KEFEA1, 2 CDK 4/6 INHIBITOR
728	1 ikn	Ω	12	162	9.00E-37	0.15	0.24		NF-KAPPA-B P65 SUBUNIT:	TRANSCRIPTION FACTOR P65: P50D:
									CHAIN: A; NF-KAPPA-B P50D	TRANSCRIPTION FACTOR, IKB/NFKB
									SUBUNIT; CHAIN: C; I-	COMPLEX
1									KAPPA-B-ALPHA; CHAIN: D;	
728	2	Ω	18	861	1.80E-32	0.27	0.23		NF-KAPPA-B P65 SUBUNIT;	TRANSCRIPTION FACTOR P65; P50D; TP ANSCRIPTION FACTOR IV BANER B
									SUBUNIT; CHAIN; C; I-	COMPLEX
									KAPPA-B-ALPHA; CHAIN: D;	!
728	լուն	m	12	162	9.00E-37	0.24	0.22		NF-KAPPA-B P65; CHAIN: A,	COMPLEX (TRANSCRIPTION
									C; NF-KAFFA-B P50; CHAIN:	REG/ANK REPEAT) COMPLEX
									B, D; I-KAPPA-B-ALPHA;	(IRANSCRIPTION REGULATION/ANK DEPEAT OF IX
728	Infi	Э	17	198	5.40E-32	0.16	0.36		NF-KAPPA-B P65- CHAIN: A	COMPLEX (TRANSCRIPTION
									C; NF-KAPPA-B P50; CHAIN:	REG/ANK REPEAT) COMPLEX
									B, D; I-KAPPA-B-ALPHA;	(TRANSCRIPTION REGULATION/ANK
									CHAIN: E, F;	REPEAT), ANKYRIN 2 REPEAT HELIX
728	Infi	យ	20	245	1.30E-26	0.03	60.0		NF-KAPPA-B P65; CHAIN: A,	COMPLEX (TRANSCRIPTION
									C; NF-KAPPA-B P50; CHAIN:	REG/ANK REPEAT) COMPLEX
									B, D; I-KAPPA-B-ALPHA;	(TRANSCRIPTION REGULATION/ANK
									CHAIN: E, F;	KEPEAI), ANKYKIN Z KEPEAI HELIX

Chain ID	, ,	Start AA 88	End AA 253	PSI- BLAST 5.40E-29	Verify score 0.06	PMF score 0.03	SeqFold	Coumpound NF-KAPPA-B P65; CHAIN: A, C, NF-KAPPA-B-80; CHAIN: B, D; I-KAPPA-B-ALPHA;	PDB annotation COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK
<	_+-+	74	208	3.90E-07	0.37	69:0		CHAIN: E, F; PHOSPHOTRIESTERASE HOMOLOGY PROTEIN; CHAIN: A, B;	REPEAT), ANKYRIN 2 REPEAT HELIX PHOSPHOTRIESTERASE PHOSPHOTRIESTERASE, HYPOTHETICAL PROTEIN
		540	618	5.40E-23	0.13	-		PNPASE; CHAIN: NULL;	SI RNA-BINDING DOMAIN POLYRIBONUCLEOTIDE NUCLEOTIDYL TRANSFERASE, SI RNA-BINDING DOMAIN, POLYNUCLEOTIDE PHOSPHORYLASE 2 (PNPASE)
4		137	217	1.10E-26	0.08	0.82		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE: CHAIN: B. C:	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
¥		361	443	7.80E-45			81.11	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B. C:	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
¥		389	469	7.80E-45	-0.04	-		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
¥		417	498	6.50E-41	-0.33	6.0		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B. C;	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
ပ		136	217	1.80E-43	90.0	96.0			COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
υ l		164	245	3.60E-46	0.27			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;	COMPLEX (ZINC FINGENDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX

				_ _	· · · · · · · · · · · · · · · · · · ·		
PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI. ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX
Coumpound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A. D.; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D, 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA: CHAIN: A, B;
SeqFold score				112.4			
PMF		0.33	_		0.95	0.92	0.19
Verify score	-0.1	0.09	0.04		-0.11	-0.04	0.01
PSI- BLAST	1.80E-50	1.40E-34	1.10E-36	2.60E-79	3.60E-38	1.60E-35	5.40E-29
End	497	282	338	444	479	499	217
Start AA	416	137	193	276	333	361	Ξ
Chain ID	O	∢	∢	¥	Y	A	S
PDB ID	Inc y	1116	1116	1116	1116	1 tf6	lubd
SEQ NO:	 	736	736	736	736	736	736

PDB annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	COMPLEY (TO ANSCRIPTION	PEGIT ATTOMONA VING VANG I	TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1;	IKANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	PECOCNITION 2 COMPET NOTEDIN	(TRANSCRIPTION REGILLATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG 1,	TRANSCRIPTION INITIATION,	INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(IRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGIT ATTONONA) VING-YANG 1:	TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-YANG I;	INTIATOR FLEMENT YYL ZINC 2	FINGER PROTEIN, DNA-PROTEIN
Coumpound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA;	CHAIN: A, B;			VVI: CHARL C. ADENO	A SCOCIATED VIBILE DE	INITIATOR ELEMENT DNA:	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;			YY1: CHAIN: C: ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA;	CHAIN: A, B;				YYI; CHAIN: C; ADENO- ASSOCIATED VIRIIS P5	INITIATOR ELEMENT DNA:	CHAIN: A, B;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	CHAIN A B.	, a , a , a , a , a , a , a , a , a , a
SeqFold score																										89.47									
PMF						-							-						-													_			
Verify		0.03				0.10	0.10						0.11						0.1													0.01			
PSI- BLAST		1.00E-56				1 300 66	1.305-33						1.20E-57						3.90E-57						2000	3.90E-57						2.60E-56			
End		273				320	272						358						385							414						469			
Start AA		162				210	017						246						274						35	304						359			
Chain ID		ນ											ن						ပ			_				 ر						ပ			
PDB ID		pqn[1.4	non.	_					Inbd						Inbd						3	pani						lubd			
SEQ ID NO:		736				71,6	000						736						736						7,12	96/		_]	736			

															_		Γ			Γ	_		
PDB annotation	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1, TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1: TRANSCRIPTION INITIATION.	INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-PROTEIN	RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG I;	INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-PROTEIN	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-FINGER GLI; GLI,	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		PROTEIN/DNA) FIVE-FINGER GLI; GLI,	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	-	PROTEIN/DNA) FIVE-FINGER GLI; GLI,	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	-	PROTEIN/DNA) FIVE-FINGER GLI; GLI,	ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	
Coumpound		YYİ; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR EI EMENT DNA-	CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS	CHAIN: A, B;			ZINC FINGER PROTEIN GLII;	CHAIN: A; DNA; CHAIN: C,	: .	ZINC FINGER PROTEIN GLII;	CHAIN: A; DNA; CHAIN: C,	Ď:	ZINC FINGER PROTEIN GLII;	CHAIN: A. DNA; CHAIN: C,	<u>~</u>	ZINC FINGER PROTEIN GLII;	CHAIN: A; DNA; CHAIN: C,	<u>``</u>	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C,
SeqFold score																			·				
PMF		-		0.96			66.0				_			-			-			_			
Verify		0		-0.18			-0.04				0.2			0.38			0.22			0.03			0.58
PSI- BLAST		1.80E-34		3.90E-51			3.60E-34				2.60E-58			1.80E-34			2.60E-70			5.20E-74			1.30E-73
End AA		469		497			497				275			300			303			359			387
Start		368	_	386			396				150	_		164			164		_	192			249
Chain ID		C		S			ပ				<			4			<			4			V
PDB ID		lubd		1ubd			1ubd				2gli			2gli			2gli			2gli	_		2gli
SEQ ID		736		736			736				736			736			736			736	_		736

Chain ID	I. I	Start AA	End	PSI- BLAST	Verify	PMF	SeqFold	Coumpound	PDB annotation
								D;	ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
<		250	387	5.20E-74			92.84	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
<		305	471	2.60E-72	-0.06	-		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
<		340	468	3.60E-33	90:00	_		ZINC FINGER PROTEIN GLI I; CHAIN: A: DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
A		360	499	2.60E-68	0.03	-		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
¥		368	496	3.60E-34	0.19	-		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BNDING PROTEIN/DNA) FIVE-FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
<		89	216	5.40E-29	-0.23	0.05		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)
L		20	230	7.20E-05			55.38	29G11 FAB; CHAIN: L, H;	CATALYTIC ANTIBODY CATALYTIC ANTIBODY, ESTERASE
∢		61	230	3.60E-05			55.09	ANTIBODY (CB 4-1); CHAIN: A, B; PEPTIDE; CHAIN: C;	COMPLEX (ANTIBODY/PEPTIDE) POLYSPECIFICITY, CROSS REACTIVITY, FAB-FRAGMENT, PEPTIDE, 2 HIV-1, COMPLEX (ANTIBODY/PEPTIDE)
٧		126	212	1.60E-17	0.19	-0.12		TELOKIN; CHAIN: A	CONTRACTILE PROTÉIN IMMUNOGLOBULIN FOLD, BETA BARREL
L L		61	230	0.0054			57.41	ANTI-ANTI-IDIOTYPE GH1002 FAB FRAGMENT; CHAIN: L. H	ANTIBODY FAB FRAGMENT ANTIBODY FAB FRAGMENT
		24	213	9.10E-18			69.87	TLYMPHOCYTE ADHESION	

INTERNATIONAL SEARCH REPORT

International application No. PCT/US01/08800

A. CLASS	SIFICATION OF SUBJECT MATTER		
IPC(7) :0	COTH 21/04; COTK 5/00; A61K 39/395; C12Q 1/68		
US CL :5	536/93.1; 530/300; 444/130.1; 435/6 5 International Patent Classification (IPC) or to both n	ational classification and IPC	
	DS SEARCHED		
	ocumentation searched (classification system followed b	oy classification symbols)	
	556/93.1; 530/500; +9+/130.1; +35/6		
searched	ion searched other than minimum documentation to t		
Electronic d	lata base consulted during the international search (nat	me of data base and, where practicable	, search terms used)
STN, med search ter	lline ms: EST, expressed sequence tags		
C. DOC	CUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appl	ropriate, of the relevant passages	Relevant to claim No.
A	VOLLRATH. D. et al. The Human Y (Map Based on Naturally Occurring Din 1992. Vol 258. pages 52-59, see whole	Deletions. Science. October	1-28
Fu	rther documents are listed in the continuation of Box C		
	Special categories of cited accuments.	"I" later document published after the int	Destion sat cline to understance
-A-	document defining the general state of the art which is not considered to be of particular relevance	the principle or theory underlying th	e In sention
	sarlies document published on or after the international filling date	"X" document of particular relevance. Il considered novel or cannot be considered.	sked to involve an inventive steb
1	the design of the second dentity of an arienty claim(s) or which is	when the document is taken alone	
	document water may involve added of another citation or other special reason (as specified)	document of particular relevance; the considered to involve an inventive step	PAPER THE GOCKMONT IN COMMENDE
-0-	document referring to an oral disclosure, use, exhibition or other means	with one or more other such docu obvious to a person skilled in the ar	l
P	document published prior to the international filing date but later than the priority date claimed	".t" document member of the same palou	
Date of t	the actual completion of the international search	Date of mailing of the international s	earch report
so SEI	PTEMBER 2001	16 NOV 2001	
Commi	nd mailing address of the ISA/US assioner of Patents and Trademarks	Authorized officer JEHANNE SOUAYA Telephone No. (703) 308-1935	1 Willi-
Box PC Washin	IT agton, D.C. 20231	TILL N (TOP) CON LOCK	/ / / /
1 5	• No. (703) 305-3930	Telephone No. (703) 308-1935 /	